FILE 'HOME' ENTERED AT 14:15:27 ON 22 FEB 2002

=> file reg COST IN U.S. DOLLARS

SINCE FILE TOTAL ENTRY SESSION 0.15 0.15

FULL ESTIMATED COST

FILE 'REGISTRY' ENTERED AT 14:15:34 ON 22 FEB 2002 USE IS SUBJECT TO THE TERMS OF YOUR STN CUSTOMER AGREEMENT. PLEASE SEE "HELP USAGETERMS" FOR DETAILS. COPYRIGHT (C) 2002 American Chemical Society (ACS)

STRUCTURE FILE UPDATES: 20 FEB 2002 HIGHEST RN 394202-19-8 DICTIONARY FILE UPDATES: 20 FEB 2002 HIGHEST RN 394202-19-8

TSCA INFORMATION NOW CURRENT THROUGH July 7, 2001

Please note that search-term pricing does apply when conducting SmartSELECT searches.

Crossover limits have been increased. See HELP CROSSOVER for details.

Calculated physical property data is now available. See HELP PROPERTIES for more information. See STNote 27, Searching Properties in the CAS Registry File, for complete details: http://www.cas.org/ONLINE/STN/STNOTES/stnotes27.pdf

The P indicator for Preparations was not generated for all of the CAS Registry Numbers that were added to the H/Z/CA/CAplus files between 12/27/01 and 1/23/02. Use of the P indicator in online and SDI searches during this period, either directly appended to a CAS Registry Number or by qualifying an L-number with /P, may have yielded incomplete results. As of 1/23/02, the situation has been resolved. Also, note that searches conducted using the PREP role indicator were not affected.

Customers running searches and/or SDIs in the H/Z/CA/CAplus files incorporating CAS Registry Numbers with the P indicator between 12/27/01 and 1/23/02, are encouraged to re-run these strategies. Contact the CAS Help Desk at 1-800-848-6533 in North America or 1-614-447-3698, worldwide, or send an e-mail to help@cas.org for further assistance or to receive a credit for any duplicate searches.

```
=> s 26062-79-3
            1 26062-79-3
                 (26062-79-3/RN)
=> d
    ANSWER 1 OF 1 REGISTRY COPYRIGHT 2002 ACS
L1
     26062-79-3 REGISTRY
RN
CN
     2-Propen-1-aminium, N,N-dimethyl-N-2-propenyl-, chloride, homopolymer
     (9CI) (CA INDEX NAME)
OTHER CA INDEX NAMES:
    Ammonium, diallyldimethyl-, chloride, polymers (8CI)
OTHER NAMES:
CN
    261LV
CN
    Additol VXT 3529
CN
    Agefloc WT 20
    Agefloc WT 20VHV
CN
CN
    Agefloc WT 2206
```

```
CN
     Agefloc WT 40
     Agefloc WT 40HV
CN
     Agefloc WT 50SLV
CN
CN
     Alcofix 109
CN
     Alcofix 182
     Aronfloc C 70
CN
     AX 04
CN
CN
     AX 05
     AX 05 (polymer)
CN
     Bufloc 536
CN
     Calgon 261
CN
     Calgon 261LV
CN
     Calgon CP 2253
CN
     Calgon CP 261XLV
CN
     Calgon CP 280
CN
CN
     Calgon DMDACC
CN
     Calgon E 904
     Calgon E 905
CN
     Calgon E 921
CN
CN
     Calgon Polymer 261
CN
     Cartafix VXT
CN
     Cat-Floc
CN
     Cat-Floc L
CN
     Cat-Floc P 112-115
     Cat-Floc T 2
CN
CN
     Cat-Floc TL
CN
     Certrex 340
CN
     CM 100
CN
     CM 100 (onium compound)
CN
     Conductive Polymer 261
CN
     CP 261
CN
     CP 261LV
CN
     CP 280
CN
     Croscolor NOFF
CN
     CV 3650
CN
     CV 3750
CN
     Danfix 707
CN
     Danfix F
CN
     Diallyldimethylammonium chloride homopolymer
     Diallyldimethylammonium chloride polymer
CN
CN
     Dimethyldiallylammonium chloride homopolymer
CN
     Dimethyldiallylammonium chloride polymer
CN
     E 261
     ECCat 2020
CN
ADDITIONAL NAMES NOT AVAILABLE IN THIS FORMAT - Use FCN, FIDE, or ALL for
     DISPLAY
     128665-21-4, 58829-44-0, 63661-21-2, 119310-15-5, 114355-07-6,
DR
37293-23-5,
     37317-00-3, 37353-76-7, 141092-78-6, 69431-41-0, 153891-18-0,
143477-08-1,
     93357-85-8, 116811-08-6, 117989-81-8, 182893-02-3, 202289-61-0,
     245064-24-8, 261769-43-1
MF
     (C8 H16 N . Cl)x
CI
     PMS, COM
PCT
     Polyvinyl
                  AGRICOLA, ANABSTR, BIOBUSINESS, BIOSIS, BIOTECHNO, CA,
     STN Files:
       CANCERLIT, CAPLUS, CEN, CHEMCATS, CHEMLIST, CIN, CSCHEM, EMBASE,
IFICDB,
       IFIPAT, IFIUDB, IPA, MEDLINE, MSDS-OHS, PIRA, PROMT, RTECS*, TOXCENTER,
       TOXLIT, USPATFULL
         (*File contains numerically searchable property data)
```

Other Sources: DSL**, TSCA** (**Enter CHEMLIST File for up-to-date regulatory information) CM1 CRN 7398-69-8 (48042-45-1) CMF C8 H16 N . Cl Me $H_2C = CH - CH_2 - N^{+} - CH_2 - CH = CH_2$ Мe ● cl -2426 REFERENCES IN FILE CA (1967 TO DATE) 108 REFERENCES TO NON-SPECIFIC DERIVATIVES IN FILE CA 2431 REFERENCES IN FILE CAPLUS (1967 TO DATE) => s diallyduhydroxypropylammonium chloride homopolymer 0 DIALLYDUHYDROXYPROPYLAMMONIUM 790501 CHLORIDE 815 CHLORIDES 790501 CHLORIDE (CHLORIDE OR CHLORIDES) 88489 HOMOPOLYMER 15 HOMOPOLYMERS 88504 HOMOPOLYMER (HOMOPOLYMER OR HOMOPOLYMERS) L20 DIALLYDUHYDROXYPROPYLAMMONIUM CHLORIDE HOMOPOLYMER (DIALLYDUHYDROXYPROPYLAMMONIUM (W) CHLORIDE (W) HOMOPOLYMER) => s diallydihydroxypropylammonium chloride homopolymer 0 DIALLYDIHYDROXYPROPYLAMMONIUM 790501 CHLORIDE 815 CHLORIDES 790501 CHLORIDE (CHLORIDE OR CHLORIDES) 88489 HOMOPOLYMER 15 HOMOPOLYMERS 88504 HOMOPOLYMER (HOMOPOLYMER OR HOMOPOLYMERS) L3 0 DIALLYDIHYDROXYPROPYLAMMONIUM CHLORIDE HOMOPOLYMER (DIALLYDIHYDROXYPROPYLAMMONIUM (W) CHLORIDE (W) HOMOPOLYMER) => file caplus medline COST IN U.S. DOLLARS SINCE FILE TOTAL ENTRY SESSION FULL ESTIMATED COST 29.10 29.25 FILE 'CAPLUS' ENTERED AT 14:22:03 ON 22 FEB 2002

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```
FILE 'MEDLINE' ENTERED AT 14:22:03 ON 22 FEB 2002
=> s l1
           2509 L1
L4
=> s obbesity or steatorrhea or hyper triglyceridemia
           1773 OBBESITY OR STEATORRHEA OR HYPER TRIGLYCERIDEMIA
=> s obbesity or steatorrhea or hypertriglyceridemia
           9278 OBBESITY OR STEATORRHEA OR HYPERTRIGLYCERIDEMIA
=> s obesity or steatorrhea or hypertriglyceridemia
          84705 OBESITY OR STEATORRHEA OR HYPERTRIGLYCERIDEMIA
=> s 17 and 14
              1 L7 AND L4
=> d ibib abs
     ANSWER 1 OF 1 CAPLUS COPYRIGHT 2002 ACS
ACCESSION NUMBER:
                           2001:63835 CAPLUS
DOCUMENT NUMBER:
                           134:131954
TITLE:
                           Fat-binding polymers for use with lipase inhibitors
INVENTOR (S):
                           Jozefiak, Thomas Henry; Mandeville, W. Harry, III;
                           Holmes-Farley, Stephen Randall; Huval, Chad Cori;
                           Garigapati, Venkata R.; Shackett, Keith K.; Concagh,
                           Danny
PATENT ASSIGNEE(S):
                           Geltex Pharmaceuticals, Inc., USA
SOURCE:
                           PCT Int. Appl., 104 pp.
                           CODEN: PIXXD2
DOCUMENT TYPE:
                           Patent
LANGUAGE:
                           English
FAMILY ACC. NUM. COUNT:
PATENT INFORMATION:
     PATENT NO.
                      KIND DATE
                                              APPLICATION NO. DATE
     -----
                              _____
                                               -----
     WO 2001005408
                        A1
                                              WO 1999-US15958 19990714
                              20010125
          W: AE, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, CA, CH, CN, CU, CZ,
              DE, DK, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS,
              JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MD, MG, MK, MN, MW, MX, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, UA, UG, UZ, VN, YU, ZA, ZW, AM, AZ, BY, KG, KZ, MD,
              RU, TJ, TM
         RW: GH, GM, KE, LS, MW, SD, SL, SZ, UG, ZW, AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, BF, BJ, CF, CG, CI, CM, GA, GN, GW, ML, MR, NE, SN, TD, TG
     AU 9949957
                         A1 20010205
                                              AU 1999-49957
                                                                  19990714
                                            WO 1999-US15958 A 19990714
PRIORITY APPLN. INFO.:
     Polymers having ether and (or) N-contg. side chains are manufd. for use in
     binding fat for treatment of obesity. A typical polymer was
     manufd. by radical polymn. of N-decylacrylamide 2.83, 3-
     acrylamidopropyltrimethylammonium chloride 18.45, and acrylamide 13.33 g.
REFERENCE COUNT:
                           10
                                  THERE ARE 10 CITED REFERENCES AVAILABLE FOR
THIS
                                  RECORD. ALL CITATIONS AVAILABLE IN THE RE
FORMAT
```

=> s therap? or pharmac? or medicin?
L9 4821358 THERAP? OR PHARMAC? OR MEDICIN?

```
=> s 19 and 14
           40 L9 AND L4
L10
=> duplicate remvove
ENTER REMOVE, IDENTIFY, ONLY, OR (?):110
'L24' IS NOT VALID HERE
Enter "REMOVE" to identify and remove duplicate answers.
Enter "IDENTIFY" to identify duplicate answers in the answer set.
Enter "ONLY" to identify and create an answer set containing only
duplicate records.
ENTER REMOVE, IDENTIFY, ONLY, OR (?):110
'L24' IS NOT VALID HERE
Enter "REMOVE" to identify and remove duplicate answers.
Enter "IDENTIFY" to identify duplicate answers in the answer set.
Enter "ONLY" to identify and create an answer set containing only
duplicate records.
ENTER REMOVE, IDENTIFY, ONLY, OR (?):?
Enter "REMOVE" to identify and remove duplicate answers.
Enter "IDENTIFY" to identify duplicate answers in the answer set.
Enter "ONLY" to identify and create an answer set containing only
duplicate records.
ENTER REMOVE, IDENTIFY, ONLY, OR (?):110
'L24' IS NOT VALID HERE
Enter "REMOVE" to identify and remove duplicate answers.
Enter "IDENTIFY" to identify duplicate answers in the answer set.
Enter "ONLY" to identify and create an answer set containing only
duplicate records.
ENTER REMOVE, IDENTIFY, ONLY, OR (?):11
'L1' IS NOT VALID HERE
Enter "REMOVE" to identify and remove duplicate answers.
Enter "IDENTIFY" to identify duplicate answers in the answer set.
Enter "ONLY" to identify and create an answer set containing only
duplicate records.
ENTER REMOVE, IDENTIFY, ONLY, OR (?):14
'L6' IS NOT VALID HERE
Enter "REMOVE" to identify and remove duplicate answers.
Enter "IDENTIFY" to identify duplicate answers in the answer set.
Enter "ONLY" to identify and create an answer set containing only
duplicate records.
ENTER REMOVE, IDENTIFY, ONLY, OR (?):15
'L9' IS NOT VALID HERE
Enter "REMOVE" to identify and remove duplicate answers.
Enter "IDENTIFY" to identify duplicate answers in the answer set.
Enter "ONLY" to identify and create an answer set containing only
duplicate records.
ENTER REMOVE, IDENTIFY, ONLY, OR (?):16
'L12' IS NOT VALID HERE
Enter "REMOVE" to identify and remove duplicate answers.
Enter "IDENTIFY" to identify duplicate answers in the answer set.
Enter "ONLY" to identify and create an answer set containing only
duplicate records.
ENTER REMOVE, IDENTIFY, ONLY, OR (?):q
'Q' IS NOT VALID HERE
Enter "REMOVE" to identify and remove duplicate answers.
Enter "IDENTIFY" to identify duplicate answers in the answer set.
Enter "ONLY" to identify and create an answer set containing only
duplicate records.
```

ENTER REMOVE, IDENTIFY, ONLY, OR (?):?

Enter "REMOVE" to identify and remove duplicate answers.

Enter "IDENTIFY" to identify duplicate answers in the answer set.

```
Enter "ONLY" to identify and create an answer set containing only
duplicate records.
ENTER REMOVE, IDENTIFY, ONLY, OR (?):
ENTER REMOVE, IDENTIFY, ONLY, OR (?):file
'FILE' IS NOT VALID HERE
Enter "REMOVE" to identify and remove duplicate answers.
Enter "IDENTIFY" to identify duplicate answers in the answer set.
Enter "ONLY" to identify and create an answer set containing only
duplicate records.
ENTER REMOVE, IDENTIFY, ONLY, OR (?):quit
'QUIT' IS NOT VALID HERE
Enter "REMOVE" to identify and remove duplicate answers.
Enter "IDENTIFY" to identify duplicate answers in the answer set.
Enter "ONLY" to identify and create an answer set containing only
duplicate records.
ENTER REMOVE, IDENTIFY, ONLY, OR (?):only
ENTER L# LIST OR (END):110
'REMVOVE' IS VALID ONLY WITH THE REMOVE OPTION
File names can only be entered when using the REMOVE option of the
DUPLICATE command.
=> d hist
     (FILE 'HOME' ENTERED AT 14:15:27 ON 22 FEB 2002)
     FILE 'REGISTRY' ENTERED AT 14:15:34 ON 22 FEB 2002
L1
              1 S 26062-79-3
              O S DIALLYDUHYDROXYPROPYLAMMONIUM CHLORIDE HOMOPOLYMER
L2
1.3
              O S DIALLYDIHYDROXYPROPYLAMMONIUM CHLORIDE HOMOPOLYMER
     FILE 'CAPLUS, MEDLINE' ENTERED AT 14:22:03 ON 22 FEB 2002
T.4
           2509 S L1
L5
           1773 S OBBESITY OR STEATORRHEA OR HYPER TRIGLYCERIDEMIA
           9278 S OBBESITY OR STEATORRHEA OR HYPERTRIGLYCERIDEMIA
L6
          84705 S OBESITY OR STEATORRHEA OR HYPERTRIGLYCERIDEMIA
L7
              1 S L7 AND L4
L8
        4821358 S THERAP? OR PHARMAC? OR MEDICIN?
L9
L10
             40 S L9 AND L4
=> d ibib abs 31-40
L10 ANSWER 31 OF 40 CAPLUS COPYRIGHT 2002 ACS
ACCESSION NUMBER:
                         1990:84026 CAPLUS
DOCUMENT NUMBER:
                         112:84026
TITLE:
                         An interpretation of the sedimentation behavior of
                         pharmaceutical kaolin and other kaolin
                         preparations in aqueous environments
                         Alexander, K. S.; Azizi, J.; Dollimore, D.; Patel, F.
AUTHOR(S):
                         Coll. Pharm., Univ. Toledo, Toledo, OH, 43606, USA
CORPORATE SOURCE:
                         Drug Dev. Ind. Pharm. (1989), 15(14-16), 2559-82
SOURCE:
                         CODEN: DDIPD8; ISSN: 0363-9045
DOCUMENT TYPE:
                         Journal
LANGUAGE:
                         English
AB
     The effect of flocculant material on kaolin-water suspensions is report
     using a pharmaceutical-grade kaolin and an industrial kaolin.
     These two systems serve as a model in a broader investigation which will
     systematically study the ancillary ingredients of pharmaceutical
     kaolin-based suspensions. These studies are based on concd. suspensions
     where fall of particles is hindered and characterized by sedimentation
"en
```

bloc" with a sludge line serving as interface between the supernatent liq.

and the settling suspension. There is an optimum concn. of macromol. flocculant which causes max. flocculation. At higher concns. of flocculant the system is stabilized. Two approaches are used to explain the results. The first represents the phenomenon as a modification of Stokes Law results. A correlation is found between the parameter A used as a characterization const. in Steinour's empirical relationship and the Richardson and Zaki exponent N. A theor. justification of this relationship is provided. A permeability relationship is 'used in the second approach based on the application of permeability equations put forward by Kozeny-Carmen. The variable k in the resultant equation is shown in theory and in practice to have a min. at some value of porosity detd. by the nature of the concd. suspension. This treatment is applied here to flocculated systems. The systems are found in general to show very high hindrance.

L10 ANSWER 32 OF 40 CAPLUS COPYRIGHT 2002 ACS

ACCESSION NUMBER: 1986:39647 CAPLUS

DOCUMENT NUMBER: 104:39647

TITLE: Preparation and performance of symplex capsules AUTHOR (S): Dautzenberg, H.; Loth, F.; Fechner, K.; Mehlis, B.;

Pommerening, K.

CORPORATE SOURCE: Inst. Polymerenchem. "Erich Correns", Ger. Acad.

Sci.,

Teltow, DDR-1530, Ger. Dem. Rep.

SOURCE: Makromol. Chem., Suppl. (1985), 9, 203-10

CODEN: MCSUEU

DOCUMENT TYPE: Journal LANGUAGE: English

Encapsulation based on the formation of polyelectrolyte complexes (symplex) was carried out by using Na cellulose sulfate (I) [9005-22-5] (polyanion) and poly(dimethyldiallylammonium chloride) (II) 26062-79-3] (polycation). I (2% aq. soln.) was treated with a 2% aq. soln. of II. The resulting capsules were washed with water, NaCl soln. or a buffer and stored until use. The kinetics of release of buserelin [57982-77-1] from the capsules was studied by using the labeled

compds. in a diffusion chamber. The effect of NaCl on the drug release was detd. A photomicrograph of the capsules formed showed that they consisted of a liq. core and a semipermeable membrane. The diam. of the capsules ranged from 0.5 to 4 mm, and the thickness of the capsules ranged

from 1 to 50 .mu.m. The capsules could be swollen to several times the original vol. by changing the osmotic pressure. The dependence of mech. strength of the capsules on the presence of NaCl in the medium was demonstrated.

L10 ANSWER 33 OF 40 CAPLUS COPYRIGHT 2002 ACS

ACCESSION NUMBER: 1985:561469 CAPLUS

DOCUMENT NUMBER: 103:161469

TITLE: Microcapsules with permeable or semipermeable walls

and a liquid core INVENTOR(S): Loth, Fritz; Dautzenberg, Horst; Pommerening, Klaus

PATENT ASSIGNEE(S): Akademie der Wissenschaften der DDR, Ger. Dem. Rep. Ger. (East), 12 pp. Addn. to Ger. (East) 160,393. SOURCE:

CODEN: GEXXA8

DOCUMENT TYPE: Patent LANGUAGE: German

FAMILY ACC. NUM. COUNT:

PATENT INFORMATION:

PATENT NO. KIND DATE APPLICATION NO. DATE ----------DD 218734 A4 19850213 DD 160393 T 19830727 DD 1981-232617 19810817 T 19830727 DD 160393 DD 1980-225200 19801114 PRIORITY APPLN. INFO.: DD 1980-225200 19801114

Capsule walls are formed by pptn. of anionic and cationic polyelectrolytes

at their interface. The anionic polyelectrolytes are sulfate or carboxylate contq. polysaccharide and/or synthetic polymers, and the cationic polyelectrolytes include quaternary ammonium surfactants and/or dyes. The microcapsules can be used for sepn. processes in preparative and anal. chem. and biochem., and in pharmacy, medicine , and agrochem. and food industries. Thus, 0.2 g Na cellulose sulfate [9005-22-5] with a degree of substitution of 0.4 was dissolved in 9.8 g H2O, and the soln. was pressed through a 0.2-mm inner diam. capillary and dropped from a height of 30 cm into a stirred bath contg. 1% aq. methylene

blue [61-73-4]. After 30 min the capsules formed were decanted and washed with H2O. The deep-blue capsules had a diam. of 3-5 mm.

L10 ANSWER 34 OF 40 CAPLUS COPYRIGHT 2002 ACS

ACCESSION NUMBER: 1975:595061 CAPLUS

83:195061 DOCUMENT NUMBER:

TITLE: Insoluble polymeric quaternary trihalogen salt coated

substrates

INVENTOR(S): Rembaum, Alan; Landel, Robert F.; Keyzer, Hendrik

PATENT ASSIGNEE(S): California Institute of Technology, USA

SOURCE:

U.S., 8 pp. CODEN: USXXAM

DOCUMENT TYPE:

Patent

LANGUAGE:

English

FAMILY ACC. NUM. COUNT:

PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE		
US 3898336	Α	19750805	US 1972-252502	19720511		
US 3778476	Α	19731211	US 1970-36431	19700511		
PRIORITY APPLN. INFO	. :		US 1970-36431	19700511		

AB Polymeric quaternary ammonium trihalides have bactericidal properties and can be deposited by pptn. on materials used in prosthesis or surgery. Thus,

trans-1,4-dichloro-2-butene-N,N,N',N'-tetramethyl-1,3-diaminopropane copolymer (I) [52193-09-6] was obtained by reacting its monomers at room temp. for 3 days in MeOH soln. I was sol. in water, but the corresponding

I3 salt was not. Other polymers prepd. included 1,4-dibromobutane-N, N, N', N'-tetramethyl-1, 3-diaminopropane copolymer [29322-33-6] and 1,3-dibromopropane-N,N,N',N'-tetramethyl-1,3-diaminopropane copolymer (II)

[29322-34-7]. Dipping Dacon cloth (which is used for arterial prosthetic devices) in aq. II soln. and then in KI-I2 soln. pptd. insol. triiodide salt of II which inhibited bacterial growth. Similarly, silk sutures and silica gel could be rendered resistant to bacteria.

L10 ANSWER 35 OF 40 MEDLINE

ACCESSION NUMBER: 97285557 MEDLINE

DOCUMENT NUMBER: 97285557 PubMed ID: 9140774

Reduction of wall adsorption in capillary zone TITLE:

electrophoresis of a basic single-chain antibody fragment

by a cationic polymeric buffer additive.

AUTHOR: Morand M; Blaas D; Kenndler E

CORPORATE SOURCE: Institute of Analytical Chemistry, University of Vienna,

Austria.

JOURNAL OF CHROMATOGRAPHY. B, BIOMEDICAL SCIENCES AND SOURCE:

APPLICATIONS, (1997 Mar 28) 691 (1) 192-6.

Journal code: CXN; 9714109. ISSN: 1387-2273.

PUB. COUNTRY: Netherlands

Journal; Article; (JOURNAL ARTICLE)

LANGUAGE: English

FILE SEGMENT: Priority Journals

ENTRY MONTH: 199707

ENTRY DATE: Entered STN: 19970805

> Last Updated on STN: 19980206 Entered Medline: 19970718

AB Reduction of adsorptive protein-wall interactions by poly(diallyldimethyl ammonium chloride), a permanently cationic polymer, at a concentration of 0.5% (w/v) is demonstrated for a basic single-chain antibody fragment (scFv, pI about 9.5) even in the range of physiological pH of around 7. The polymer additive forms a positively charged layer at the silica surface which reverses electroosmosis and leads to electrostatic repulsion

of the positively charged basic protein.

L10 ANSWER 36 OF 40 MEDITNE

ACCESSION NUMBER: 96091252 MEDLINE

DOCUMENT NUMBER: 96091252 PubMed ID: 7489112

TITLE: [The antimutagenic activity of ternary diallyl

copolymers].

Antimutagennaia aktivnost' troinykh sopolimerov

diallil'nogo riada.

AUTHOR: Aleksandrova V A; Kotliarova E B; Odin A P; Domnina N S;

Shevchenko V A; Topchiev D A

SOURCE: RADIATSIONNAIA BIOLOGIIA, RADIOECOLOGIIA, (1995 Sep-Oct)

35

(5) 746-51.

Journal code: BWZ; 9317212. ISSN: 0869-8031.

PUB. COUNTRY: RUSSIA: Russian Federation

Journal; Article; (JOURNAL ARTICLE)

LANGUAGE: Russian

FILE SEGMENT: Priority Journals

ENTRY MONTH: 199601

ENTRY DATE: Entered STN: 19960125

> Last Updated on STN: 19960125 Entered Medline: 19960102

AB Antimutagenic activity of triple copolymers of diallyl origin was investigated by animal cell test (mouse bone marrow erythrocytes, 1.5 Gy of gamma irradiation) and by plant cell test (seeds of barley, 5 Gy of gamma irradiation). Effective protection of genetic structure was achieved

owing to combination of moderate antimutagenic activity of the polymer matrix and scavenging ability of sterically hindered phenols in the polymer side chain.

L10 ANSWER 37 OF 40 MEDLINE

ACCESSION NUMBER: 88101911 MEDLINE

DOCUMENT NUMBER: 88101911 PubMed ID: 2962394

TITLE: [Capsular mosaicism and characteristics of the adsorptive

interaction of Treponema pallidum in vitro].

Izuchenie kapsuliarnoi mozaichnosti i osobennostei adsorbtsionnogo vzaimodeistviia blednykh treponem in

vitro.

AUTHOR: Milich M V; Skripkin Iu K; Fedorova D L; Topchiev D A;

Bednova V N

SOURCE: VESTNIK DERMATOLOGII I VENEROLOGII, (1987) (9) 28-33.

Journal code: X9U; 0414246. ISSN: 0042-4609.

PUB. COUNTRY:

Journal; Article; (JOURNAL ARTICLE)

LANGUAGE: Russian

FILE SEGMENT: Priority Journals

ENTRY MONTH: 198802

ENTRY DATE: Entered STN: 19900305

> Last Updated on STN: 19900305 Entered Medline: 19880210

L10 ANSWER 38 OF 40 MEDLINE

ACCESSION NUMBER: 85284418 MEDLINE

PubMed ID: 3839743 DOCUMENT NUMBER: 85284418

TITLE: [Chemiluminescence of peritoneal macrophages activated by

non-natural polyelectrolytes].

Khemiliuminestsentsiia peritoneal'nykh makrofagov, aktivirovannaia neprirodnymi polielektrolitami.

AUTHOR: Korkina L G; Suslova T B; Guliaeva Zh G; Zezin A B;

Velichkovskii B T

SOURCE: DOKLADY AKADEMII NAUK SSSR, (1985 May-Jun) 282 (1) 206-9.

Journal code: EBK; 7505465. ISSN: 0002-3264.

PUB. COUNTRY: USSR

Journal; Article; (JOURNAL ARTICLE)

LANGUAGE: Russian

FILE SEGMENT: Priority Journals

ENTRY MONTH: 198510

ENTRY DATE: Entered STN: 19900320

> Last Updated on STN: 19900320 Entered Medline: 19851003

L10 ANSWER 39 OF 40 MEDLINE

ACCESSION NUMBER: 82061521 MEDLINE

DOCUMENT NUMBER: 82061521 PubMed ID: 7302387

Effect of three bile acid binding polymers on the TITLE:

biosynthesis of 14C-cholesterol from 14C-sodium acetate in

the rat.

AUTHOR: Gilfillan J L; Huff J W

SOURCE: RESEARCH COMMUNICATIONS IN CHEMICAL PATHOLOGY AND

PHARMACOLOGY, (1981 Aug) 33 (2) 373-6.

Journal code: R62; 0244734. ISSN: 0034-5164.

PUB. COUNTRY: United States

Journal; Article; (JOURNAL ARTICLE)

LANGUAGE: English

FILE SEGMENT: Priority Journals

ENTRY MONTH: 198201

ENTRY DATE: Entered STN: 19900316

> Last Updated on STN: 19900316 Entered Medline: 19820109

AB The relative activity of three bile acid binding polymers in increasing cholesterol biosynthesis in the rat from 14C-acetate was determined by measuring blood levels of 14C-cholesterol after intraperitoneally administered 14C-acetate. CAT-FLOC and 3,3-ionene were 4-5 times more active than cholestyramine in this study which correlated well with the results of hypocholesteremic testing in dogs.

L10 ANSWER 40 OF 40 MEDLINE

ACCESSION NUMBER: 81109797 MEDLINE DOCUMENT NUMBER: 81109797 PubMed ID: 7193034

TITLE: The bile acid binding and hypocholesterolemic action of

two

water-soluble polymers.

AUTHOR: Kuron G W; Grier N; Huff J W

SOURCE: ATHEROSCLEROSIS, (1980 Nov) 37 (3) 353-60.

Journal code: 95X; 0242543. ISSN: 0021-9150.

PUB. COUNTRY: Netherlands

Journal; Article; (JOURNAL ARTICLE)

LANGUAGE: English

FILE SEGMENT: Priority Journals

ENTRY MONTH: 198103

ENTRY DATE: Entered STN: 19900316

> Last Updated on STN: 19900316 Entered Medline: 19810317

AB The in vitro bile acid binding properties of 2 water-soluble, linear, cationic resins, poly-[(dimethylimino)trimethylene chloride] or 3,3-ione C1, and poly-diallyldimethylammonium chloride) or CAT-FLOC were determined. Both polymers were substantially more active than cholestyramine. All were compared for hypocholesterolemic effect in normo-cholesterolemic dogs. CAT-FLOC and 3,3-ionene C1, administered at 1.8 and 1.2 g/day, respectively, exhibited cholesterol-lowering action equivalent to cholesteryramine given at 12 g/day. The results of this study suggest that effective reduction of plasma cholesterol may be achieved with significantly lower doses of bile acid sequestrants.

=> d ibib abs 21-30

L10 ANSWER 21 OF 40 CAPLUS COPYRIGHT 2002 ACS

ACCESSION NUMBER: 1995:973665 CAPLUS

DOCUMENT NUMBER: 124:4489

TITLE: Ion-capture reagents and methods for performing

binding assays

INVENTOR(S): Hiltibran, Robert G.; Jou, Yi-Her; Stroupe, Stephen

D.; Kline, Steven J.; Schultz, Steven G.

PATENT ASSIGNEE(S): Abbott Laboratories, USA SOURCE:

PCT Int. Appl., 87 pp.

CODEN: PIXXD2

DOCUMENT TYPE: Patent LANGUAGE: English

FAMILY ACC. NUM. COUNT:

PATENT INFORMATION:

APPLICATION NO. DATE DATE PATENT NO. KIND DATE ----------WO 1995-US3168 19950314 WO 9525282 A1 19950921

W: AU, CA, JP

RW: AT, BE, CH, DE, DK, ES, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE A1 19951003 AU 1995-19978 19950314 AU 9519978 PRIORITY APPLN. INFO.: US 1994-214017 19940315 WO 1995-US3168 19950314

AB This invention presents novel reagents, sepn. techniques and assay procedures, esp. immunoassay procedures, which allow both the indicator and the capture reagents to be in soln. to avoid problems of slowed reaction kinetics. The sepn. procedure involves a sol. capture reagent, comprising a specific binding member attached to a charged substance, and an insol. solid phase that is oppositely charged with respect to the charged substance included in the capture reagent. A test sample suspected of contg. the analyte of interest, e.g., antigen, drug,

hormone,

is mixed with the capture reagent to form a charged capture reagent/analyte complex. The reaction mixt. is contacted to the oppositely charged solid phase to attract, attach, and sep. the capture reagent/analyte complexes and any unreacted capture reagent from the reaction mixt. With an appropriate indicator reagent, i.e., a second specific binding member which is conjugated to a label capable of producing a detectable signal, both sandwich and competitive assays can

be

performed.

L10 ANSWER 22 OF 40 CAPLUS COPYRIGHT 2002 ACS

ACCESSION NUMBER: 1995:267191 CAPLUS

DOCUMENT NUMBER: 122:59438

TITLE: Modified metal oxide layer as support for active

materials and reagents

INVENTOR(S): Boettcher, Horst; Kallies, Karl-Heinz

PATENT ASSIGNEE(S): Germany

SOURCE: Ger. Offen., 14 pp.

CODEN: GWXXBX

DOCUMENT TYPE: Patent LANGUAGE: German

FAMILY ACC. NUM. COUNT: 1

PATENT INFORMATION:

PATENT NO. KIND DATE APPLICATION NO. DATE

DE 4308146 A1 19940922 DE 1993-4308146 19930315

AB Metal oxide layers (Al2O3, SiO3, TiO2) are treated with penetrating agents

(salts, orgs., polymers) during formation from gases (CVD, PVD) or solns.(sol-gel process), to increase the porosity of the layer or to change the structure of the layer to increase the absorptivity for active materials. The layers are useful in cosmetics, chem. anal., medical diagnosis, pharmaceuticals.

L10 ANSWER 23 OF 40 CAPLUS COPYRIGHT 2002 ACS

ACCESSION NUMBER: 1995:92522 CAPLUS

DOCUMENT NUMBER: 122:38691

TITLE: Synthesis and biological activity of polymer salts of

benzylpenicillin based on cationic polyelectrolytes

containing polydialkyldiallylammonium

AUTHOR(S): Aleksandrova, V. A.; Zlobina, V. A.; Dmitriyev, G.

Α.;

Milonova, T. I.; Fedorova, D. L.; Topchiyev, D. A.

CORPORATE SOURCE: Inst. Neftekhim. Sint., Moscow, Russia SOURCE: Khim.-Farm. Zh. (1994), 28(5), 38-40

CODEN: KHFZAN; ISSN: 0023-1134

DOCUMENT TYPE: Journal LANGUAGE: Russian

AB Benzylpenicillin was immobilized on bioactive polymers contg. diallyl group and antibacterial activity of prodrugs obtained was evaluated. The in vitro sensitivity of the resistant strain Staphylococcus aureus to the drug was enhanced possible due to the membrane-active properties of the polymer matrix. In vivo, the drug was tested in rabbits infected with Treponema pallidum. The effects of the drug were evident after 48 h. Thus, a combination of the bactericidal properties of the polymer matrix and benzylpenicillin is suitable for the treatment of syphilis requiring

reduced amt. of antibiotic for therapeutic effects.

L10 ANSWER 24 OF 40 CAPLUS COPYRIGHT 2002 ACS

ACCESSION NUMBER:

1993:503026 CAPLUS

DOCUMENT NUMBER:

119:103026

TITLE:

Dentinal desensitizing compositions containing

polyelectrolytes

INVENTOR (S):

Lim, Richard M.; Herms, James Keeth; Fertel, William

S.; Synodis, Joseph

PATENT ASSIGNEE(S): SOURCE:

Block Drug Co. Inc., USA Eur. Pat. Appl., 10 pp.

CODEN: EPXXDW

DOCUMENT TYPE:

Patent

LANGUAGE:

English

FAMILY ACC. NUM. COUNT:

PATENT INFORMATION:

I	PATENT NO.				KII	ND	DATE		APPLICATION NO.					ο.	DATE			
-																		
I	EP 549281			A1		19930630			EP 1992-311594				4	19921218				
I	EΡ	EP 549281			В:	l	19990414											
		R:	ΑT,	BE,	CH,	DE,	DK,	ES,	FR,	GB,	GR,	ΙE,	IT,	LI,	LU,	MC,	NL,	PT,
SE																		
τ	US	5270	031		Α		1993	1214		US	199	91-83	1181	1	1991	1220		
1	ΑU	9230	141		A:	l	1993	0624		AU	199	92-30	141		1992	1214		
1	ΑU	6556	95		B	2	1995	0105										
I	BR	9204	695		Α		1993	0622		BR	199	92-46	595		1992	1217		
I	ΑT	1787	86		E		1999	0415		AT	199	92-3	1159	4	1992	1218		
I	ES	2133	309		T3	3	1999	0916		ES	199	92-33	1159	4	1992	1218		
(CA	2085	975		A/	A	1993	0621		CA	199	92-20	0859	75	1992	1221		
(CA	2085	975		C		1999	0330										
j	JP	0606	5083		A2	2	1994	0308		JP	199	92-3	5633	8	1992	1221		
ن	JP	2821	967		B2	2	1998	1105										
PRIOR	ITY	APP	LN.	INFO.	:				υ	JS 19	91-8	3118	11		1991	1220		

AB Polyelectrolytes contg. a functional group of carboxylate, carboxy, amino alkylammonium or mixts. thereof are included in a dentifrice base or other

oral compns. for relieving pain and discomfort caused by hypersensitive teeth. For example, a dentifrice contg. Me vinyl ether-maleic acid copolymer K salt was formulated.

L10 ANSWER 25 OF 40 CAPLUS COPYRIGHT 2002 ACS

ACCESSION NUMBER:

1992:113518 CAPLUS

DOCUMENT NUMBER:

116:113518

TITLE:

Bile acid remover-containing pharmaceutical

spheres for lowering blood cholesterol

PATENT ASSIGNEE(S):

Eureka, Inc., USA

SOURCE:

Jpn. Kokai Tokkyo Koho, 7 pp.

CODEN: JKXXAF

DOCUMENT TYPE:

Patent

LANGUAGE:

Japanese

FAMILY ACC. NUM. COUNT:

PATENT INFORMATION:

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE									
	JP 03170436	A2	19910724	JP 1989-307375	19891127									
AB	The title drugs,	based	on removal of	bile acids from c	irculation to									
	promote conversi	on of	chlorestol to	bile acids, are ma	nufd. by (1)									
	dissolving nondegradable polymers (e.g. polystyrene sulfonate-													
	vinylbenzyltrime	thylam	monium chlorid	e copolymer) in a										
	stirring a bile	acid r	emover (e.g. 1	,5-dimethyl-1,5-										
	diazoundecamethy	lenepo	lymethyl bromi	de) with the soln.	or give a									
				persion with a sol										

 $% \left(1\right) =\left(1\right) +\left(1\right) +\left($

that is nonmiscible with the bile and remover) to prep. the spheres for oral administration. The side effects are minimal.

L10 ANSWER 26 OF 40 CAPLUS COPYRIGHT 2002 ACS

ACCESSION NUMBER: 1991:542256 CAPLUS

DOCUMENT NUMBER: 115:142256

TITLE: Pharmaceutical compositions containing HMG

CoA reductase inhibitor and/or squalene synthetase inhibitor for treating peripheral atherosclerotic

disease

INVENTOR(S): Eisman, Martin

PATENT ASSIGNEE(S): Squibb, E. R., and Sons, Inc., USA

SOURCE: Eur. Pat. Appl., 17 pp.

CODEN: EPXXDW

DOCUMENT TYPE: Patent LANGUAGE: English

FAMILY ACC. NUM. COUNT: 1

PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
EP 401705	A2	19901212	EP 1990-110475	19900601
EP 401705	A3	19930107		
R: AT, BE,	CH, DE	, DK, ES, FR,	GB, GR, IT, LI, LU	, NL, SE
CA 2016467	AA	19901205	CA 1990-2016467	19900510
AU 9054950	A1	19901206	AU 1990-54950	19900511
HU 54059	A2	19910128	HU 1990-3320	19900604
JP 03020226	A2	19910129	JP 1990-147164	19900605
ZA 9004310	Α	19910327	ZA 1990-4310	19900605
PRIORITY APPLN. INFO	. :		US 1989-361520	19890605
OTHER SOURCE(S):	MA	RPAT 115:1422	56	

AB 3-Hydroxy-3-methylglutaryl (HMG) CoA reductase and/or squalene synthetase inhibitors are used to prep. pharmaceuticals for treating arteriosclerosis obliterans and/or intermittent claudication in mammals. Optionally, a pharmaceutical is included which reduces serum cholesterol by a mechanism other than inhibiting prodn. of HMG CoA reductase or squalene synthetase. Tablet and capsule formulations are given. One contained pravastatin (HMG CoA reductase inhibitor) 7, lactose

67, microcryst. cellulose 20, croscarmellose Na 2, Mg stearate 1, and Mg oxide 3 parts by wt.

L10 ANSWER 27 OF 40 CAPLUS COPYRIGHT 2002 ACS

ACCESSION NUMBER: 1991:538744 CAPLUS

DOCUMENT NUMBER: 115:138744

TITLE: Asbestos-free filter for treatment of cationic and

anionic suspensions

INVENTOR(S): Oertel, Ulrich; Geyer, Stefan; Petzold, Gudrun;

Buchhamer, Heide Marie; Schwarz, Simona

PATENT ASSIGNEE(S): Akademie der Wissenschaften der DDR, Fed. Rep. Ger.

SOURCE: Ger. (East), 3 pp.

CODEN: GEXXA8

DOCUMENT TYPE: Patent

LANGUAGE: German

FAMILY ACC. NUM. COUNT: 1

PATENT INFORMATION:

PATENT NO. KIND DATE APPLICATION NO. DATE

19910508 DD 289711 A5 DD 1989-335252 19891205

AB The filter comprises a cationic and an anionic layer with thickness ratio 9:1 to 1:1. The filter is suitable for treatment of food and medicine for sterilization or ultrapurifn. A suitable filter comprises a kieselguhr-cellulose filter support coated with poly(dimethyldiallylammonium chloride) on 1 side and a polyacrylate latex on the other side.

L10 ANSWER 28 OF 40 CAPLUS COPYRIGHT 2002 ACS

ACCESSION NUMBER: 1991:516915 CAPLUS

DOCUMENT NUMBER:

115:116915

TITLE:

Preparation of asbestos-free filters for treatment of

cationic and anionic suspensions

INVENTOR(S):

Oertel, Ulrich; Geyer, Stefan

PATENT ASSIGNEE(S):

Akademie der Wissenschaften der DDR, Fed. Rep. Ger.

SOURCE:

Ger. (East), 3 pp. CODEN: GEXXA8

DOCUMENT TYPE:

Patent

LANGUAGE:

German

FAMILY ACC. NUM. COUNT:

PATENT INFORMATION:

PATENT NO. KIND DATE APPLICATION NO. DATE DD 289710 A5 19910508 DD 1989-335251 19891205

The filters comprise an asbestos-free layer having pos. .zeta.-potential, AΒ where .gtoreq.1 of the constituents of the layer contains a cationic deposit or coating of org. polycations and low-mol. wt. amphiphilic org. anions. Optionally, the org. anions contain an acid group and have 12-13 C atoms. The filter is suitable for treatment of food and medicine for sterilization or ultrapurifn. A suitable filter comprises kieselguhr and milled spruce cellulose coated with an aq. soln. of poly(dimethyldiallyl ammonium chloride) and Na stearate (optionally Na palmitate or Na abietate).

L10 ANSWER 29 OF 40 CAPLUS COPYRIGHT 2002 ACS

ACCESSION NUMBER:

1991:478972 CAPLUS

DOCUMENT NUMBER:

115:78972

TITLE:

Method for impregnation of drugs on textiles

INVENTOR (S):

Ryl'tsev, V. V.; Vasil'eva, T. S.; Filatov, V. N.; Kabanov, V. A.; Zezin, A. B.; Rogacheva, V. B.; Oltarzhevskaya, N. D.; Krichevskii, G. E.; Subbotko,

O. A.; et al.

PATENT ASSIGNEE(S):

All-Union Scientific-Research Institute of the Textile-Haberdashery Industry, USSR; Moscow State

SOURCE:

U.S.S.R. From: Otkrytiya, Izobret. 1990, (48), 112.

CODEN: URXXAF

DOCUMENT TYPE:

Patent

LANGUAGE:

Russian

FAMILY ACC. NUM. COUNT:

PATENT INFORMATION:

KIND DATE APPLICATION NO. DATE PATENT NO. 19901230 SU 1988-4444332 19880428 -----A1 SU 1617069

AB A procedure for ensuring medicinal properties of textile material made of cellulose and polyamide fibers by applying a compn. contg. biocompatible polymer, drug, and H2O to a jersey material of 1 .times. 1 rib weave by printing with subsequent drying is improved. The drug content in the material and air permeability and moisture absorption of the material are increased by using a blend of polyacrylic acid with 3.6 .times. 103-105 mol. wt. and polydimethyldiallylammonium chloride (I) with 7.5 .times. 103-106 mol. wt. (1:2 wt. ratio) as the biocompatible polymer. Thus, a compn. contained a blend of polyacrylic acid with 3.6 .times. 103-105 mol. wt. and I with 7.5 .times. 103-106 mol. wt. (1:2 wt. ratio) 10-40, drug 1-6, NaCl 3-5 wt. %, and the balance being H2O.

L10 ANSWER 30 OF 40 CAPLUS COPYRIGHT 2002 ACS

ACCESSION NUMBER: 1991:88411 CAPLUS

DOCUMENT NUMBER: 114:88411

TITLE: Liquid compositions containing crystallizable

compounds and cationic polymers

INVENTOR (S):

Nishida, Yuichi

PATENT ASSIGNEE(S):

Lion Corp., Japan

SOURCE:

Jpn. Kokai Tokkyo Koho, 11 pp.

CODEN: JKXXAF

DOCUMENT TYPE:

Patent

LANGUAGE:

Japanese

FAMILY ACC. NUM. COUNT:

PATENT INFORMATION:

PATENT NO. KIND APPLICATION NO. DATE DATE ------JP 02166161 A2 19900626 JP 1988-320073 19881219

GI

AΒ A liq. compn. having a wide industrial application contains .gtoreq.1 EtOH-sol. cationic polymer dissolved in an org. solvent and a crystallizable compd. such as fatty acid deriv. The cationic polymer may be I (R1, R2 = H, C1-3 alkyl, Ph; R3, R4 = C1-18 alkyl, H; Y = anion; s, t

Ι

= 0, 1; s + t = 1; m .gtoreq. 1 and n .gtoreq. 0 to produce polymer with mol. wt. 10,000-1.5 .times. 106), diallylmethylammonium chloride-2-hydroxyethyl cellulose copolymer, etc. The lig. compn. in water or in org. solvents is stable at low temp. and has only little ppts.

A nonionic surfactant with <10 HLB may be added. Thus, a compn. consisted

of dimethyldiallylammonium chloride polymer 0.5, octanoic acid glyceride 1.0, EtOH 91.5, and POE hydrogenated castor oil 7% by wt. This compn. was

stable at least 50 days at low temp. The pptn. of the fatty acid deriv. was prevented by the presence of the cationic polymer. A hair tonic contg. pentadecanoic acid glyceride 3, Merckote-100 0.1, citric acid 0.3, 1-menthol 0.1, sorbitan monooleate 0.8, a perfume 0.5, and EtOH to 100.0% was prepd.

```
=> s polyelectrolyte
        23558 POLYELECTROLYTE
=> d hist
     (FILE 'HOME' ENTERED AT 14:15:27 ON 22 FEB 2002)
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L1
             1 S 26062-79-3
L2
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L3
              0 S DIALLYDIHYDROXYPROPYLAMMONIUM CHLORIDE HOMOPOLYMER
     FILE 'CAPLUS, MEDLINE' ENTERED AT 14:22:03 ON 22 FEB 2002
L4
          2509 S L1
          1773 S OBBESITY OR STEATORRHEA OR HYPER TRIGLYCERIDEMIA
L<sub>5</sub>
          9278 S OBBESITY OR STEATORRHEA OR HYPERTRIGLYCERIDEMIA
L6
T.7
         84705 S OBESITY OR STEATORRHEA OR HYPERTRIGLYCERIDEMIA
L8
             1 S L7 AND L4
        4821358 S THERAP? OR PHARMAC? OR MEDICIN?
L9
            40 S L9 AND L4
L10
          23558 S POLYELECTROLYTE
T.11
=> 111 and 17
L11 IS NOT A RECOGNIZED COMMAND
The previous command name entered was not recognized by the system.
For a list of commands available to you in the current file, enter
"HELP COMMANDS" at an arrow prompt (=>).
=> s 111 and 17
L12
            1 L11 AND L7
=> d ibib abs
L12 ANSWER 1 OF 1 CAPLUS COPYRIGHT 2002 ACS
ACCESSION NUMBER: 2000:686244 CAPLUS DOCUMENT NUMBER: 133:256576
TITLE:
                       Slimming sprays containing a hydrosoluble
film-forming
                        polymer
INVENTOR(S):
                        Picard, Elisabeth
PATENT ASSIGNEE(S):
                        L'oreal, Fr.
SOURCE:
                        Eur. Pat. Appl., 20 pp.
                        CODEN: EPXXDW
DOCUMENT TYPE:
                        Patent
LANGUAGE:
                        French
FAMILY ACC. NUM. COUNT: 1
PATENT INFORMATION:
    PATENT NO.
                 KIND DATE
                                        APPLICATION NO. DATE
     -----
     EP 1038518
                     A1 20000927
                                          EP 2000-400531 20000228
        R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT,
            IE, SI, LT, LV, FI, RO
                A1 20000929
     FR 2791258
                                          FR 1999-3532
                                                          19990322
     FR 2791258
                      B1 20010824
PRIORITY APPLN. INFO.:
                                       FR 1999-3532
                                                        A 19990322
    Slimming sprays contg. a C1-6 monoalc., a hydrosol. film-forming polymer
     and slimming active ingredient are disclosed. A slimming sprays
    Lviset CA66 (vinyl acetate-crotonic acid copolymer) 4.5, aminomethyl
```

propanol 0.48, Cola nitida ext. 0.25, dried ext. of guarana fruit 0.1, ext. of Coleus barbatus contg. 60% forskolin 0.1, and ethanol 94.57%.

REFERENCE COUNT: 6 THERE ARE 6 CITED REFERENCES AVAILABLE FOR THIS

RECORD. ALL CITATIONS AVAILABLE IN THE RE

FORMAT

=> d ibib abs 110 1-20

L10 ANSWER 1 OF 40 CAPLUS COPYRIGHT 2002 ACS ACCESSION NUMBER: 2001:178354 CAPLUS

DOCUMENT NUMBER: 134:212761

TITLE: Controlled, phased-release suppository and its method

of production

SOURCE: U.S., 10 pp.
CODEN: USXXAM

DOCUMENT TYPE: Patent LANGUAGE: English

FAMILY ACC. NUM. COUNT: 1

PATENT INFORMATION:

PATENT NO. KIND DATE APPLICATION NO. DATE

US 6200590 B1 20010313 US 1998-131591 19980810

AB A phased-release suppository delivery system is disclosed wherein microscopic polymeric "nanospheres" ladened with one or more active agents

are homogeneously incorporated within a pharmaceutically acceptable suppository base. The prepn. of the "nanospheres" allows the spheres to be transported, substantially intact, across fenestrated membranes such as the capillary membranes of the rectum. The method of prepn. of the "nanospheres" allows for the controlled release of active agent(s) only after a substantial no. of the spheres have been transported

across the capillary membrane of the rectum or other body cavity and have been taken up into the systemic circulation system.

REFERENCE COUNT:

THERE ARE 6 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE

FORMAT

L10 ANSWER 2 OF 40 CAPLUS COPYRIGHT 2002 ACS ACCESSION NUMBER: 2000:772736 CAPLUS

6

DOCUMENT NUMBER: 133:305579

TITLE: Artificial cells microencapsulated genetically

engineered E. coli DH5 cells for the removal of

undesired electrolytes and/or metabolites

INVENTOR(S): Prakash, Satya; Chang, Thomas M. S.

PATENT ASSIGNEE(S): McGill University, Can. SOURCE: PCT Int. Appl., 34 pp.

CODEN: PIXXD2

DOCUMENT TYPE: Patent LANGUAGE: English

FAMILY ACC. NUM. COUNT: 1

PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE		
WO 2000065030	A2	20001102	WO 2000-CA482	20000427		
WO 2000065030	A3	20010125				

W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, CA, CH, CN, CR, CU, CZ, DE, DK, DM, DZ, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, TZ, UA, UG, US, UZ, VN, YU, ZA, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM

RW: GH, GM, KE, LS, MW, SD, SL, SZ, TZ, UG, ZW, AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, BF, BJ, CF,

CG, CI, CM, GA, GN, GW, ML, MR, NE, SN, TD, TG

PRIORITY APPLN. INFO.: US 1999-131468 P

The present invention relates to a compn. for the removal of at least one undesired electrolyte and/or metabolite in a patient, which comprises a genetically engineered E. coli DH5 cells microencapsulated in artificial cells to be capable of removing said undesired electrolyte and/or metabolite, wherein said undesired electrolyte is selected from the group consisting of K, Mg, P, Na, Cl and said undesired metabolite is selected from the group consisting of uric acid, cholesterol, bilirubin, and creatinine, wherein said removal of undesired electrolyte and/or metabolite lowers the undesired chem. concn. to a therapeutically acceptable level.

L10 ANSWER 3 OF 40 CAPLUS COPYRIGHT 2002 ACS ACCESSION NUMBER: 2000:654904 CAPLUS

DOCUMENT NUMBER: 133:213101

TITLE: Poly(dimethyldiallylammonium chloride) for treatment

of periodontitis

INVENTOR (S): Klimov, V. I.; Zaklyuchaeva, V. I.; Boyarkina, N. M.;

Kuznetsova, L. I.; Leshchankina, E. L.

PATENT ASSIGNEE(S): Zakrytoe Aktsionernoe Obshchestvo Firma "TOKEM",

Russia

Patent

SOURCE: Russ. From: Izobreteniya 1999, (13), 376.

CODEN: RUXXE7

DOCUMENT TYPE:

LANGUAGE:

Russian

FAMILY ACC. NUM. COUNT:

PATENT INFORMATION:

PATENT NO. KIND DATE APPLICATION NO. DATE ----------RU 2129857 C1 19990510 RU 1997-102782 19970224

AB Title only translated.

L10 ANSWER 4 OF 40 CAPLUS COPYRIGHT 2002 ACS

ACCESSION NUMBER: 2000:573840 CAPLUS

DOCUMENT NUMBER: 133:179157

TITLE: Derivatized microfibrillar polysaccharides, their

formation and use in dispersions

INVENTOR (S): Cash, Mary Jean; Chan, Anita N.; Conner, Herbert

Thompson; Cowan, Patrick Joseph; Gelman, Robert Alan; Lusvardi, Kate Marritt; Thompson, Samuel Anthony;

Tise, Frank Peine

PATENT ASSIGNEE(S): Hercules Incorporated, USA

SOURCE: PCT Int. Appl., 84 pp.

CODEN: PIXXD2

DOCUMENT TYPE: Patent

LANGUAGE: English

FAMILY ACC. NUM. COUNT:

PATENT INFORMATION:

KIND DATE PATENT NO. APPLICATION NO. DATE -----

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WO 2000047628
                      A2
                            20000817
                                          WO 2000-US3319
                                                           20000208
                            20001207
     WO 2000047628
                      A3
         W: AE, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, CA, CH, CN, CU, CZ,
            DE, DK, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS,
             JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MD, MG, MK,
            MN, MW, MX, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ,
             TM, TR, TT, UA, UG, UZ, VN, YU, ZA, ZW, AM, AZ, BY, KG, KZ, MD,
             RU, TJ, TM
         RW: GH, GM, KE, LS, MW, SD, SL, SZ, TZ, UG, ZW, AT, BE, CH, CY, DE,
             DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, BF, BJ, CF,
             CG, CI, CM, GA, GN, GW, ML, MR, NE, SN, TD, TG
     BR 2000005116
                     Α
                          20010102
                                         BR 2000-5116
                          20010228
                                         EP 2000-911740
                      A2
                                                           20000208
            AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT,
             IE, SI, LT, LV, FI, RO
     NO 2000005085
                     Α
                           20001207
                                          NO 2000-5085
                                                           20001009
PRIORITY APPLN. INFO.:
                                        US 1999-248246
                                                        A 19990210
                                        WO 2000-US3319
                                                        W 20000208
     The invention is directed to the followings. A method for producing
     derivatized microfibrillar polysaccharide, including but not limited to
     cellulose, derivatized by steric and/or electrostatic forces, where the
     electrostatic forces are provided by anionic charge or by a combination
of
     both anionic and cationic charge, by stabilizing and/or microfibrillating
     a polysaccharide starting material. A method of modifying the rheol.
     properties of a compn. of matter using derivatized microfibrillar
    polysaccharide. Method of improving coatings, paper manuf., and the
     stability of emulsions, dispersions, and foams using a derivatized
     microfibrillar polysaccharide. Compns. that include derivatized
     microfibrillar polysaccharide, e.g., CM cellulose, including paper
     compns., comestible compns., non-comestible spreadable compns.
     (cosmetics), and emulsions, dispersion, and foams.
L10 ANSWER 5 OF 40 CAPLUS COPYRIGHT 2002 ACS
ACCESSION NUMBER:
                         2000:68383 CAPLUS
DOCUMENT NUMBER:
                         132:113118
TITLE:
                         Polyelectrolyte coatings on biological templates
INVENTOR(S):
                        Neu, Bjoern; Baeumler, Hans; Donath, Edwin; Moya,
                         Sergio; Sukhorukov, Gleb; Moehwald, Helmuth; Caruso,
                         Frank
PATENT ASSIGNEE(S):
                        Max-Planck-Gesellschaft zur Foerderung der
                         Wissenschaften e.V., Germany
SOURCE:
                         PCT Int. Appl., 43 pp.
                         CODEN: PIXXD2
DOCUMENT TYPE:
                         Patent
LANGUAGE:
                         German
FAMILY ACC. NUM. COUNT:
PATENT INFORMATION:
     PATENT NO.
                     KIND DATE
                                          APPLICATION NO. DATE
                     ----
     -----
                                           ------------
     WO 2000003797
                      A1
                            20000127
                                          WO 1999-EP5063
                                                           19990715
         W: JP, US
        RW: AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL,
            PT, SE
                                                           19980715
    EP 972563
                           20000119
                                          EP 1998-113181
        R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT,
            IE, SI, LT, LV, FI, RO
    DE 19907552
                      A1
                           20000831
                                          DE 1999-19907552 19990222
    EP 1098696
                      A1
                           20010516
                                          EP 1999-938268 19990715
```

R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT,

IE, FI

PRIORITY APPLN. INFO.:

EP 1998-113181 A 19980715 DE 1999-19907552 A 19990222 WO 1999-EP5063 W 19990715

Capsules provided with a polyelectrolyte covering are produced by use of AΒ template particles comprising biol. cells, cell aggregates, subcellular structures, virus particles, aggregates of biomols. such as immune complexes, or aggregates of amphiphilic materials such as liposomes or micelles. These template particles are coated with successive layers of oppositely charged polyelectrolytes. The template particles are subsequently lysed, leaving empty capsule shells (microcapsules) which

may

be loaded with active agents such as enzymes, drugs, polymers, dyes, sensor mols., agrochems., or flavorings. The permeability of the microcapsules can be regulated by varying the conditions of polyelectrolyte deposition or by incorporation of surfactants and/or lipids. Polymn. reactions may be carried out within the microcapsules,

as

the capsules are impermeable to the polymeric reaction products, but permeable to monomers; the course of polymn. can be controlled through control of capsule permeability and external conditions (e.g. medium compn.) in ways not possible with bulk polymn. Thus, washed erythrocytes were fixed in 25% aq. glutardialdehyde and coated alternately with pos.-charged poly(allylamine)-HCl (50-60 kDa, 0.5 q/dL) and neq.-charged Na polystyrenesulfonate (70 kDa, 0.5 g/dL) by adsorption; the coating steps were repeated 5 times. The erythrocyte cores of the capsules were removed by lysis with 1.2% NaOCl soln.; their removal was signaled by a decrease in turbidity.

REFERENCE COUNT:

THERE ARE 4 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE

FORMAT

L10 ANSWER 6 OF 40 CAPLUS COPYRIGHT 2002 ACS

ACCESSION NUMBER:

1999:613767 CAPLUS

DOCUMENT NUMBER:

131:247144

TITLE:

Fabrication of multilayer-coated particles and hollow

shells by electrostatic self-assembly of

nanocomposite

multilayers on decomposable colloidal templates INVENTOR(S): Caruso, Frank; Caruso, Rachel Anne; Donath, Edwin;

Mohwald, Helmuth; Sukhorukov, Gleb

PATENT ASSIGNEE(S):

Max-Planck-Gesellschaft Zur Forderung Der

Wissenschaften E.V., Germany

SOURCE:

PCT Int. Appl., 55 pp.

CODEN: PIXXD2

DOCUMENT TYPE:

Patent

LANGUAGE:

English

FAMILY ACC. NUM. COUNT:

PATENT INFORMATION:

PATENT NO.	KIND DATE	APPLICATION NO.	DATE
WO 9947253	A1 19990923	WO 1999-EP1854	19990319
W: JP, US			
RW: AT, BE,	CH, CY, DE, DK, ES,	FI, FR, GB, GR, IE,	IT, LU, MC, NL,
PT, SE			
DE 19812083	A1 19990930	DE 1998-19812083	19980319
EP 972563	A1 20000119	EP 1998-113181	19980715
R: AT, BE,	CH, DE, DK, ES, FR,	GB, GR, IT, LI, LU,	NL, SE, MC, PT,
IE, SI,	LT, LV, FI, RO		
EP 1064088	A1 20010103	EP 1999-916840	19990319

R: AT, BE, CH, DE, DK, FR, GB, IT, LI, NL, SE PRIORITY APPLN. INFO.: DE 1998-19812083 A 19980319 EP 1998-113181 A 19980715

WO 1999-EP1854 W 19990319

Coated particles and hollow shells (av. diam. .ltoreq.15 .mu.m) are AB produced by coating colloidal particles, e.g., polystyrene templates, with

alternating layers of oppositely charged nanoparticles and polyelectrolytes, then optionally removing the colloidal templates or cores. The nanoparticles can be inorg., e.g., SiO2, or org., including biomols., e.g., proteins. The template can be removed by disintegration using thermal, chem. or pH treatment, e.g., calcination or decompn. upon exposure to solvents or low pH. The wall thickness of the hollow spheres can be controlled by varying the no. of nanoparticle deposition cycles, and the size and shape are detd. by the morphol. of the templating colloid. The shells may contain an active agent, such as pharmaceuticals, herbicides, pesticides, catalysts, pigments. Applications can include slow or targeted release of the active substances.

REFERENCE COUNT:

THERE ARE 2 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE

FORMAT

L10 ANSWER 7 OF 40 CAPLUS COPYRIGHT 2002 ACS

ACCESSION NUMBER: DOCUMENT NUMBER:

1999:473475 CAPLUS 132:69232

TITLE:

Development of cellulose sulfate-based

polyelectrolyte

complex microcapsules for medical applications Dautzenberg, Horst; Schuldt, Ute; Grasnick, Gerd; Karle, Peter; Muller, Petra; Lohr, Matthias;

Pelegrin,

AUTHOR (S):

Mireia; Piechaczyk, Marc; Rombs, Kerstin V.;

Gunzburg,

Walter H.; Salmons, Brian; Saller, Robert M.

University of Potsdam, Potsdam, Germany

Ann. N. Y. Acad. Sci. (1999), 875 (Bioartificial

SOURCE: Organs

II), 46-63

CODEN: ANYAA9; ISSN: 0077-8923

PUBLISHER:

New York Academy of Sciences

DOCUMENT TYPE:

CORPORATE SOURCE:

Journal

LANGUAGE:

English

Microencapsulation, as a tool for immunoisolation for allogenic or xenogenic implants, is a rapidly growing field. However most of the approaches are based on alginate/polylysine capsules, despite this system's obvious disadvantages such as its pyrogenicity. The authors report a different encapsulation system based on sodium cellulose sulfate and polydiallyldimethyl ammonium chloride for the encapsulation of mammalian cells. The authors have characterized this system regarding capsule formation, strength and size of the capsules as well as viability of the cells after encapsulation. In addn., the authors demonstrate the efficacy of these capsules as a "microfactory" in vitro and in vivo. Using encapsulated hybridoma cells the authors were able to demonstrate long-term release of antibodies up to four months in vivo. In another application the authors could show the therapeutic relevance of encapsulated genetically modified cells as an in vivo activation center for cytostatic drugs during tumor therapy.

REFERENCE COUNT:

33 THERE ARE 33 CITED REFERENCES AVAILABLE FOR

THIS

RECORD. ALL CITATIONS AVAILABLE IN THE RE

L10 ANSWER 8 OF 40 CAPLUS COPYRIGHT 2002 ACS ACCESSION NUMBER: 1999:113552 CAPLUS DOCUMENT NUMBER: 130:173009 TITLE: Combinations of HMG-CoA reductase inhibitors and nicotinic acid and methods for treating hyperlipidemia INVENTOR(S): Bova, David J.; Dunne, Josephine Kos Pharmaceuticals, Inc., USA PATENT ASSIGNEE(S): PCT Int. Appl., 86 pp. SOURCE: CODEN: PIXXD2 DOCUMENT TYPE: Patent English LANGUAGE: FAMILY ACC. NUM. COUNT: PATENT INFORMATION: PATENT NO. KIND DATE APPLICATION NO. DATE A1 19990211 WO 1998-US15989 19980731 ------WO 9906046 W: AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, CA, CH, CN, CU, CZ, DE, DK, EE, ES, FI, GB, GE, GH, GM, HR, HU, ID, IL, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MD, MG, MK, MN, MW, MX, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, UA, UG, UZ, VN, YU, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM RW: GH, GM, KE, LS, MW, SD, SZ, UG, ZW, AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, BF, BJ, CF, CG, CI, CM, GA, GN, GW, ML, MR, NE, SN, TD, TG US 2001006644 A1 20010705 US 1997-903871 19970731 AU 9886800 A1 19990222 AU 1998-86800 19980731 EP 1003515 A1 20000531 EP 1998-938227 19980731 R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE, FI NO 2000000407 20000316 NO 2000-407 20000127 PRIORITY APPLN. INFO.: US 1997-903871 A 19970731 WO 1998-US15989 W 19980731 The present invention relates to solid pharmaceutical AB combinations for oral administration comprising nicotinic acid or a nicotinic acid compd. or mixts. thereof in an extended release form and an HMG-CoA reductase inhibitor, which are useful for altering lipid levels in subjects suffering from, for example, hyperlipidemia and atherosclerosis, without causing drug-induced hepatotoxicity, myopathy or rhabdomyolysis. The present invention also relates to methods of altering serum lipids in subjects to treat, for example, hyperlipidemia in hyperlipidemics, lipidemia in normolipidemics diagnosed with or predisposed to cardiovascular disease, and atherosclerosis, by administering such oral solid pharmaceutical combinations once per day as a single dose during the evening hours, without causing drug-induced hepatotoxicity, myopathy or rhabdomyolysis, or without causing in at least an appreciable no. of individuals drug-induced hepatotoxicity, myopathy or rhabdomyolysis to such a level that discontinuation of such therapy would be required. More particularly, the present invention concerns oral solid

required. More particularly, the present invention concerns oral solid pharmaceutical combinations comprised of, for example, (1) an HMG-CoA reductase inhibitor for immediate or extended release, (2) nicotinic acid, a nicotinic acid compd. or mixts. thereof, and (3) a swelling agent to form a sustained release compn. for extended release of the nicotinic acid or nicotinic acid compd. or mixts. thereof for nocturnal or evening dosing for reducing serum lipids and increasing HDL-cholesterol. In accordance with the present invention, and by way of

example, a compn. for oral administration during the evening hours to alter serum lipids comprised of nicotinic acid and hydroxypropyl Me cellulose in the form of an extended or sustained release tablet or caplet

coated with a coating comprising an HMG-CoA reductase inhibitor in immediate release form is disclosed. Also in accordance with the present invention, the pharmaceutical combinations may include a nonsteroidal anti-inflammatory agent for reducing the capacity of nicotinic acid or nicotinic acid compds. to provoke flushing reactions in individuals.

REFERENCE COUNT:

11

THERE ARE 11 CITED REFERENCES AVAILABLE FOR

THIS

RECORD. ALL CITATIONS AVAILABLE IN THE RE

FORMAT

L10 ANSWER 9 OF 40 CAPLUS COPYRIGHT 2002 ACS

ACCESSION NUMBER:

1999:81698 CAPLUS

DOCUMENT NUMBER:

130:144187

TITLE:

Immobilization of vitamin A acid by cationic

polyelectrolytes

INVENTOR(S):

Thuenemann, Andreas

PATENT ASSIGNEE(S):

Max-Planck-Gesellschaft zur Foerderung der

Wissenschaften e.V., Germany

SOURCE:

Ger. Offen., 22 pp.

CODEN: GWXXBX

DOCUMENT TYPE:

Patent

LANGUAGE:

German

FAMILY ACC. NUM. COUNT:

PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO. DATE
DE 19732139	A1	19990128	DE 1997-19732139 19970725
WO 9904821	A2	19990204	WO 1998-EP4644 19980724
WO 9904821	A3	19990415	
W: CA, JP,	US		
RW: AT, BE,	CH, CY	, DE, DK,	ES, FI, FR, GB, GR, IE, IT, LU, MC, NL,
PT, SE			
EP 1003559	A2	20000531	EP 1998-942616 19980724
R: AT, BE,	CH, DE	, DK, ES,	FR, GB, GR, IT, LI, LU, NL, SE, MC, PT,
IE, FI			
JP 2001510809	T2	20010807	JP 2000-503872 19980724
PRIORITY APPLN. INFO	. :		DE 1997-19732139 A 19970725
			WO 1998-EP4644 W 19980724

Amphiphilic vitamin A acid is immobilized and thereby stabilized in AB pharmaceutical dosage forms by formation of mesomorphic complexes with cationic polyelectrolytes. These complexes are sol. in polar org. solvents such as MeOH, EtOH, 2-BuOH, iso-PrOH, and CHCl3; they dissoc. in these solvents, but remain assocd. in solvents of lower polarity. The complexes form lamellar viscoelastic films which can be used e.g. for treatment of skin diseases or as part of a photosynthetic system.

0.5% aq. soln. of poly(diallyldimethylammonium chloride) (mol. wt. 180,000) was added dropwise to a soln. of vitamin A acid in aq. NaOH until

no further pptn. occurred. The pptd. complex was dissolved in MeOH and purified by ultrafiltration to remove excess vitamin A acid and NaCl. The

complex formed an optically anisotropic film.

L10 ANSWER 10 OF 40 CAPLUS COPYRIGHT 2002 ACS

ACCESSION NUMBER:

1998:732910 CAPLUS

DOCUMENT NUMBER:

CORPORATE SOURCE:

130:16558

TITLE:

Nanoengineering of inorganic and hybrid hollow

spheres

by colloidal templating

AUTHOR (S):

Caruso, Frank; Caruso, Rachel A.; Mohnwald, Helmuth

Max Planck Inst. Colloids Interfaces, Berlin,

D-72489,

SOURCE:

Science (Washington, D. C.) (1998), 282(5391),

1111-1114

CODEN: SCIEAS; ISSN: 0036-8075

PUBLISHER:

American Association for the Advancement of Science

DOCUMENT TYPE:

Journal English

LANGUAGE:

Hollow silica and silica-polymer spheres with diams. 720-1000 nm were fabricated by consecutively assembling silica nanoparticles and polymer onto colloids and subsequently removing the templated colloid either by calcination or decompn. upon exposure to solvents. SEM and TEM images demonstrate that the wall thickness of the hollow spheres can be readily controlled by varying the no. of nanoparticle-polymer deposition cycles,

and the size and shape are detd. by the morphol. of the templating colloid. The hollow spheres produced are envisioned to have applications in areas ranging from medicine to pharmaceutics to

materials science.

REFERENCE COUNT:

29 THERE ARE 29 CITED REFERENCES AVAILABLE FOR

THIS

RECORD. ALL CITATIONS AVAILABLE IN THE RE

FORMAT

L10 ANSWER 11 OF 40 CAPLUS COPYRIGHT 2002 ACS

ACCESSION NUMBER:

1998:539993 CAPLUS

DOCUMENT NUMBER:

129:341317

TITLE:

Recent progress in research on immobilization of

microorganisms and animal cells on microcapsules

AUTHOR (S):

Mei, Lehe; Yao, Shanjing

CORPORATE SOURCE:

Department of Chemical Engineering, Zhejiang University, Hangzhou, 310027, Peop. Rep. China

SOURCE:

Xiandai Huagong (1998), 18(1), 19-22

CODEN: HTKUDJ; ISSN: 0253-4320

PUBLISHER:

Zhongguo Huagong Xinxi Zhongxin

DOCUMENT TYPE:

Journal; General Review

LANGUAGE:

Chinese

A review with 12 refs. discussing research on the immobilization of microorganisms or animal cells on microcapsules with emphasis on characteristics and application of NaCS-PDMDAAC biomicrocapsules in fermn., food and drug manufg., biochem anal., clin. diagnosis, sustained-release dosage forms and others.

L10 ANSWER 12 OF 40 CAPLUS COPYRIGHT 2002 ACS

ACCESSION NUMBER:

1998:527187 CAPLUS

DOCUMENT NUMBER:

129:166216

TITLE:

Ophthalmic compositions including glycerin and

propylene glycol

INVENTOR(S):

Hu, Zhenze; Denick, John

PATENT ASSIGNEE(S):

Bausch & Lomb Incorporated, USA PCT Int. Appl., 16 pp.

SOURCE:

CODEN: PIXXD2

Patent

DOCUMENT TYPE:

LANGUAGE:

English

FAMILY ACC. NUM. COUNT:

PATENT INFORMATION:

```
APPLICATION NO. DATE
    PATENT NO.
                   KIND DATE
    WO 9832421 A1 19980730 WO 1998-US1649 19980129
        W: AL, AU, BB, BR, BY, CA, CU, EE, ES, FI, GB, HU, IL, JP, KE, KG,
            KP, KR, MD, MK, MN, MW, NO, NZ, PL, PT, RU, SE, SI, SK, TJ, TR,
            TT, UA, UZ, VN
        RW: GH, GM, KE, LS, MW, SD, SZ, UG, ZW, AT, BE, CH, DE, DK, ES, FI,
            FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, BF, BJ, CF, CG, CI, CM,
            GA, GN, ML, MR, NE, SN, TD, TG
                   A 19980901
                                       US 1997-794690
                                                       19970129
    AU 9862533
                    A1 19980818
                                       AU 1998-62533
                                                       19980129
    AU 723024
                    B2 20000817
    EP 969812
                    A1 20000112
                                       EP 1998-904734
                                                       19980129
        R: DE, ES, FR, GB, IT, IE
                                       BR 1998-7117
    BR 9807117 A 20000425
                                                      19980129
    JP 2001511135
                     T2 20010807
                                       JP 1998-532253
                                                       19980129
                                     US 1997-794690 A 19970129
PRIORITY APPLN. INFO.:
                                     WO 1998-US1649 W 19980129
    There are disclosed ophthalmic compns. having high water-binding
    properties which are useful as: moisturizing and lubricating (i.e.
```

AB There are disclosed ophthalmic compns. having high water-binding properties which are useful as: moisturizing and lubricating (i.e. artificial tear) solns., dry eye therapies, contact lens wetting and lubricating solns., and as delivery vehicles for ophthalmic drugs. The subject compns. include glycerin in combination with propylene glycol.

The subject compns. may further include cellulose derivs., e.g. hydroxypropyl Me cellulose, along with preservatives, e.g. benzylalkonium chloride, PHMB, sorbic acid, etc.. Preferred compns. have at least 11 % bound water, a pH from about 7.1 to 7.5, and an osmolality between about 280 to about 320 mOsm/Kg. A moisturizing eye drop formulation contained glycerol 1.0, propylene glycol 0.5, HPMC 1.0, boric acid 0.300, Na borate 0.035, NaCl 0.096, KCl 0.097, EDTA 0.030, benzalkonium chloride (50%) 0.021, a purified water to 100% wt./wt.

L10 ANSWER 13 OF 40 CAPLUS COPYRIGHT 2002 ACS

ACCESSION NUMBER: 1998:479401 CAPLUS

DOCUMENT NUMBER: 12

129:113593

TITLE:

Encapsulated cells producing antibodies

INVENTOR(S):

Piechaczyk, Marc; Pelegrin, Mireia; Marin, Mariana;

Saller, Robert; Salmons, Brian

PATENT ASSIGNEE(S):

Bavarian Nordic Research Institute A/S, Den.

SOURCE:

PCT Int. Appl., 22 pp.

CODEN: PIXXD2

DOCUMENT TYPE:

Patent

LANGUAGE:

English

FAMILY ACC. NUM. COUNT:

PATENT INFORMATION:

P	ATE	ENT I	NO.		KI	ND	DATE APPLICATION NO.								DATE				
-										-									
WO 9827966 A2				2	1998	0702		W	0 19	97-E	P712	0	19971218						
WO 9827966 A3				3	19981112														
		W:	AL,	AM,	ΑT,	ΑU,	ΑZ,	BA,	BB,	BG,	BR,	BY,	CA,	CH,	CN,	CU,	CZ,	DE,	
			DK,	EE,	ES,	FI,	GB,	GE,	GH,	GM,	HU,	ID,	IL,	IS,	JP,	ΚE,	KG,	KΡ,	
			KR,	ΚZ,	LC,	LK,	LR,	LS,	LT,	LU,	LV,	MD,	MG,	MK,	MN,	MW,	MX,	NO,	
			NZ,	PL,	PT,	RO,	RU,	SD,	SE,	SG,	SI,	SK,	SL,	TJ,	TM,	TR,	TT,	UA,	
			ŪĠ,	US,	UΖ,	VN,	YU,	ZW,	AM,	ΑZ,	BY,	KG,	ΚZ,	MD,	RU,	ΤJ,	TM		
		RW:	GH,	GM,	ΚE,	LS,	MW,	SD,	SZ,	UG,	ZW,	ΑT,	BE,	CH,	DE,	DK,	ES,	FI,	
			FR,	GB,	GR,	ΙE,	IT,	LU,	MC,	NL,	PT,	SE,	BF,	ВJ,	CF,	CG,	CI,	CM,	
			GA,	GN,	ML,	MR,	NE,	SN,	TD,	TG									

AU 9862059 A1 19980717 AU 1998-62059 19971218 EP 948319 A2 19991013 EP 1997-954822 19971218

R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT,

IE, SI, LT, LV, FI, RO

JP 2001507344 T2 20010605 JP 1998-528353 19971218 PRIORITY APPLN. INFO.: DK 1996-1497 A 19961223 WO 1997-EP7120 W 19971218

AB The present invention relates to capsules encapsulating

antibody-producing

cells and to the use of such capsules and encapsulated

cells, and to the use of such capsules and encapsulated cells, resp., for implantation in vivo for long term delivery or sustained delivery of antibodies of **therapeutic** interest.

L10 ANSWER 14 OF 40 CAPLUS COPYRIGHT 2002 ACS

ACCESSION NUMBER: 1997:682782 CAPLUS

DOCUMENT NUMBER: 127:336527

TITLE: Immobilization of Retinoic Acid by Cationic

Polyelectrolytes Thuenemann, Andreas

CORPORATE SOURCE: Max Planck Institut fuer Kolloid-

Grenzflaechenforschung, Teltow-Seehof, D-14513,

Germany

SOURCE: Langmuir (1997), 13(23), 6040-6046

CODEN: LANGD5; ISSN: 0743-7463

PUBLISHER: American Chemical Society

DOCUMENT TYPE: Journal LANGUAGE: English

AB Retinoic acid was immobilized by pptg. its complexes with cationic polyelectrolytes from aq. soln. Polyelectrolytes with different architectures, such as poly(ionene-6,3 bromide), poly(dimethyldiallylammonium chloride), and poly(N-methyl-4-vinylpyridinium chloride), form self-assembling complexes contg. retinoic acid (70% (wt./wt.)). All these complexes are thermodynamically stable and can be processed into mesomorphously ordered films with interesting phys. properties. In contrast to the brittle cryst. retinoic acid the complexes with polyelectrolytes are highly deformable viscoelastic materials. All materials show lamellar mesophase structures; their Tq

value strongly depends on the polyelectrolyte. It is suggested that these

materials have great potential as **pharmaceutical** agents as well as models for the investigation and the mimicking of chromophores in visual pigments and photosynthetic bacteria. The properties of the complexes are examd. by X-ray diffraction, DSC, polarization optical microscopy, UV-vis spectroscopy, and stress-strain measurements.

L10 ANSWER 15 OF 40 CAPLUS COPYRIGHT 2002 ACS ACCESSION NUMBER: 1997:513569 CAPLUS

DOCUMENT NUMBER: 127:181203

TITLE: Microencapsulated genetically engineered

microorganisms for clinical application

INVENTOR(S): Chang, Thomas M. S.; Prakash, Satya

PATENT ASSIGNEE(S): McGill University, Can.; Chang, Thomas M. S.;

Prakash,

AUTHOR(S):

Satya

SOURCE: PCT Int. Appl., 36 pp.

CODEN: PIXXD2

DOCUMENT TYPE: Patent LANGUAGE: English

FAMILY ACC. NUM. COUNT: 1

PATENT INFORMATION:

```
APPLICATION NO. DATE
                   KIND DATE
    PATENT NO.
     -----
                                        _____
    WO 9726903 A1 19970731 WO 1997-CA40
                                                        19970120
        W: CA, JP, US
        RW: AT, BE, CH, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT,
SE
                     A1 19981111
                                        EP 1997-900520 19970120
        R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT,
            IE, FI
                      T2 19990309
                                        JP 1997-526371 19970120
    JP 11502875
    JP 3228941
                    B2 20011112
    US 6217859
                    B1 20010417
                                        US 1998-117099 19980722
PRIORITY APPLN. INFO.:
                                      GB 1996-1333 A 19960123
                                      WO 1997-CA40
                                                     W 19970120
    The present invention relates to a compn. for oral administration to a
AB
    patient for the removal of undesired chems. and/or amino acids caused by
а
    disease, which comprises a microorganism entrapped or microencapsulated
to
    be capable of removing the undesired chems. and/or amino acids in assocn.
    with a pharmaceutically acceptable carrier for oral
    administration to the patient.
L10 ANSWER 16 OF 40 CAPLUS COPYRIGHT 2002 ACS
                      1997:463766 CAPLUS
ACCESSION NUMBER:
DOCUMENT NUMBER:
                       127:126433
                       Comparative in vitro cytotoxicity studies of
TITLE:
                       polycations for gene therapy
AUTHOR (S):
                       Fischer, D.; Zange, R.; Kissel, T.
CORPORATE SOURCE:
                       Dept. of Pharmaceutical Technology and
                       Biopharmaceutics, Marburg, 35032, Germany
                        Proc. Int. Symp. Controlled Release Bioact. Mater.
SOURCE:
                        (1997), 24th, 647-648
                        CODEN: PCRMEY; ISSN: 1022-0178
PUBLISHER:
                        Controlled Release Society, Inc.
DOCUMENT TYPE:
                        Journal
LANGUAGE:
                        English
    The cytotoxicity in L929 mouse fibroblasts was assessed for polycations
    considered for use as DNA-condensing agents for gene therapy.
    Data are presented for poly-DADMAC, polyvinyl pyridinium bromide, human
    serum albumin, DEAE-dextran, poly-L-lysine, polyethylenimine, and human
    serum albumin cationized by coupling to hexamethylenediamine.
L10 ANSWER 17 OF 40 CAPLUS COPYRIGHT 2002 ACS
ACCESSION NUMBER:
                        1997:179549 CAPLUS
DOCUMENT NUMBER:
                        126:324659
TITLE:
                        Poly(diallyldimethylammonium chloride) as a cationic
                        coating for capillary electrophoresis
                       Liu, Qicai; Lin, Fangming; Hartwick, Richard A.
AUTHOR(S):
CORPORATE SOURCE:
                        Paul M. Gross Chemical Lab., Duke University, Durham,
                       NC, 27708, USA
SOURCE:
                        J. Chromatogr. Sci. (1997), 35(3), 126-130
                        CODEN: JCHSBZ; ISSN: 0021-9665
PUBLISHER:
                        Preston Publications
DOCUMENT TYPE:
                        Journal
LANGUAGE:
                        English
    A novel cationic polymer coating that exhibits a fast anodal
    electroosmotic flow (EOF) was developed for capillary electrophoresis.
Ιn
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the 1st approach, poly(diallyldimethylammonium chloride) is chem. bonded onto the interior capillary wall; in a 2nd approach, the polymer is phys.

adsorbed onto the inner wall of the capillary. Capillaries modified by both approaches exhibit an anodal EOF in the pH range of 2.2-8.8, with a relatively pH-independent EOF (.apprx.-5.5 .times. 10-4 cm2/V s) over the pH range of 2.2-5.5. The application of the novel EOF-reversed phase is demonstrated by the improved sepn. of basic proteins and beta-adrenergic blocking drugs. Sepn. efficiencies ranging from 50,000 to 200,000 plates per m are obsd. for proteins. The relative std. deviation of migration times for multiple injections of test proteins is <0.65%. The reproducibility of capillary synthesis is 2.3% relative std. deviation

for

capillaries synthesized on three different days. The lifetimes of both the bonded and phys. coated capillaries exceed 40 h of continuous use at 240 V/cm at pH 4.

L10 ANSWER 18 OF 40 CAPLUS COPYRIGHT 2002 ACS

ACCESSION NUMBER: 1996:666260 CAPLUS

DOCUMENT NUMBER: 125:338842

TITLE: Study of the inter-polyelectrolyte reactions between

cellulose graft copolymers containing acidic groups

and antimicrobial polycations

AUTHOR(S): Virnik, A. D.; Penenzhik, M. A.; Ryshkina, I. S.;

Kozhanova, T. Ya.; Zezi, A. B.; Rogacheva, V. B.

CORPORATE SOURCE: "A. N. Kosygin" Moscow State Textile Academy, Moscow,

117918, Russia

SOURCE: Cellul. Chem. Technol. (1996), 30(1-2), 39-47

CODEN: CECTAH; ISSN: 0576-9787

DOCUMENT TYPE: Journal

LANGUAGE: English

AB Interpolymer reactions involving antimicrobial polycations (polyhexamethylene guanidine and polydimethyldiallylammonium) and differently structured cellulose graft copolymers contg. acidic groups were studied in order to det. the principles of developing antimicrobial cellulose fibrous materials with controlled properties. It was found

that

the kinetics of the interpolymer reaction, the compn. and properties of the resulting polyelectrolyte complexes depend heavily on the structure of

antimicrobial polycation and that of cellulose graft polyanion.

L10 ANSWER 19 OF 40 CAPLUS COPYRIGHT 2002 ACS

ACCESSION NUMBER: 1996:392656 CAPLUS

DOCUMENT NUMBER: 125:95894

TITLE: Immobilization of biological matter by

polyelectrolyte

complex formation

AUTHOR(S): Dautzenberg, H.; Lukanoff, B.; Eckert, U.; Tiersch,

B.; Schuldt, U.

CORPORATE SOURCE: Univ. Potsdam, Germany

SOURCE: Ber. Bunsen-Ges. (1996), 100(6), 1045-1053

CODEN: BBPCAX; ISSN: 0940-483X

DOCUMENT TYPE: Journal LANGUAGE: English

AB The immobilization of biol. matter plays an important part in creating biol. active systems for applications preferably in biotechnol. processes or in medical treatments. The immobilization by polyelectrolyte complex formation, allowing the encapsulation of sensitive biol. objects or materials without causing any damage to them, has become a powerful alternative method. In more detail exptl. results on capsule formation with sodium cellulose sulfate (NaCS) and poly(diallyldimethylammonium chloride) as reaction compds. are reported. Capsule formation and capsule

properties (morphol. features, cut-off, rate of NaCS conversion, size and mech. stability) are looked at under polymer-chem. and physicochem. aspects. The importance of osmotic effects is revealed. It is shown

that

the capsule properties depend not only on the chem. structure of the polyelectrolytes or the conditions of capsule prepn., but also remarkably on the polymer-chem. characteristics of the reaction components, particularly on their mol. mass and/or mol. mass distribution.

L10 ANSWER 20 OF 40 CAPLUS COPYRIGHT 2002 ACS ACCESSION NUMBER: 1995:997220 CAPLUS

DOCUMENT NUMBER: 124:127123

TITLE: Metal oxide composite with controllable release of

bioactive agents

INVENTOR(S): Boettcher, Horst; Kallies, Karl-Heinz; Marx, Joerg

PATENT ASSIGNEE(S): Feinchemie GmbH Sebnitz, Germany

SOURCE: Eur. Pat. Appl., 21 pp.

CODEN: EPXXDW

DOCUMENT TYPE: Patent LANGUAGE: German

FAMILY ACC. NUM. COUNT: 1

PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO. DATE
EP 680753	A2	19951108	EP 1995-106926 19950508
EP 680753	A3	19960828	
R: DE, ES,	FR, GB	, IT	
DE 4416003	A1	19951109	DE 1994-4416003 19940506
DE 4416003	C2	19990211	
DE 4416001	A1	19951130	DE 1994-4416001 19940506
DE 4416001	C2	19971211	
PRIORITY APPLN. INFO	.:		DE 1994-4416001 19940506
		,	DE 1994-4416003 19940506

AB A controlled-release dosage form of a bioactive agent comprises a metal oxide matrix, the homogeneously dispersed active agent, and .gtoreq.1 substance which regulates the release of the active agent. Among the latter are low-mol.-wt. water-sol. substances, polyionic compds., microporous fillers, high-boiling solvents, and penetration enhancers. The composite is useful for prepn. of sustained-release formulations of drugs, cosmetics, bactericides, insecticides, and pesticides. Thus, an SiO2 sol was prepd. by mixing (EtO)4Si 50, EtOH 200, and 0.01N HCl 100 mL and stirring for 20 h. Carbamazepine (1 g) was dissolved in a corresponding amt. of sol, with or without addn. of 1.0 g sorbitol, 3 mL poly(dimethyldiallylammonium chloride), or 2 g di-Bu phthalate. The sol was then neutralized to pH 7, which induced gelation, and the gel was dried and granulated. Poly(dimethyldiallylammonium chloride) slowed the release of carbamazepine from the granules.

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L4

(FILE 'HOME' ENTERED AT 14:15:27 ON 22 FEB 2002)

FILE 'REGISTRY' ENTERED AT 14:15:34 ON 22 FEB 2002

L1 1 S 26062-79-3

L2 0 S DIALLYDUHYDROXYPROPYLAMMONIUM CHLORIDE HOMOPOLYMER
L3 0 S DIALLYDIHYDROXYPROPYLAMMONIUM CHLORIDE HOMOPOLYMER

FILE 'CAPLUS, MEDLINE' ENTERED AT 14:22:03 ON 22 FEB 2002 2509 S L1

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L5
          1773 S OBBESITY OR STEATORRHEA OR HYPER TRIGLYCERIDEMIA
L6
          9278 S OBBESITY OR STEATORRHEA OR HYPERTRIGLYCERIDEMIA
L7
          84705 S OBESITY OR STEATORRHEA OR HYPERTRIGLYCERIDEMIA
L8
             1 S L7 AND L4
L9
       4821358 S THERAP? OR PHARMAC? OR MEDICIN?
L10
            40 S L9 AND L4
L11
         23558 S POLYELECTROLYTE
L12
             1 S L11 AND L7
=> s 14 and dihydroxyprop?
            2 L4 AND DIHYDROXYPROP?
=> s 113 not 18
            1 L13 NOT L8
=> d ibib abs
L14 ANSWER 1 OF 1 CAPLUS COPYRIGHT 2002 ACS
ACCESSION NUMBER:
                       1994:279841 CAPLUS
DOCUMENT NUMBER:
                        120:279841
TITLE:
                        Cationic cellulose derivatives containing fatty
                        quaternium groups in a pre-shampoo conditioning
                        composition
INVENTOR (S):
                        Tashjian, Anne
PATENT ASSIGNEE(S):
                        USA
SOURCE:
                        U.S., 7 pp.
                        CODEN: USXXAM
DOCUMENT TYPE:
                        Patent
LANGUAGE:
                        English
FAMILY ACC. NUM. COUNT:
PATENT INFORMATION:
                    KIND DATE
     PATENT NO.
                                         APPLICATION NO. DATE
     -----
                                         -----
                                         US 1992-884964 19920515
                     Α
                          19940222
    US 5288484
AB
    A pretreatment conditioner comprising (1) an aq. system of .apprx.0.1-20%
     of a cationic cellulose deriv. quaternized with fatty C10-C18 alkyl
     groups, (2) .apprx.0.05-20% of a quaternary polymer, and (3)
     .apprx.0.02-10% of a quaternary ammonium salt. A shampoo conditioner
     contained hydroxyethyl cellulose 0.5, polyquaternium-6 2.0, dimethicone
     copolyol 2.5, propylene glycol 4.0, behenamidopropyl
     dihydroxypropyl PG dimonium chloride 1.0, Crodacel QS 4.0, Na
     citrate 0.04, fragrances, solubilizers, preservatives, dyes, botanicals
     5.124, and water q.s. 100%.
=> s diallyldihydroxylpropylammonium
            0 DIALLYLDIHYDROXYLPROPYLAMMONIUM
=> s diallyl dihydroxylpropyl ammonium
            O DIALLYL DIHYDROXYLPROPYL AMMONIUM
=> s n,n-diallyl-n-methyl-n-(2,3-dihydroxypropyl) ammonium
MISSING OPERATOR '-METHYL-N-(2,3-DIHYDR'
The search profile that was entered contains terms or
nested terms that are not separated by a logical operator.
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=> s 321904-07-8

0 321904-07-8

=> s 321904-08-9 1 321904-08-9 => d ibib abs L18 ANSWER 1 OF 1 CAPLUS COPYRIGHT 2002 ACS ACCESSION NUMBER: 2001:63835 CAPLUS DOCUMENT NUMBER: 134:131954 TITLE: Fat-binding polymers for use with lipase inhibitors INVENTOR (S): Jozefiak, Thomas Henry; Mandeville, W. Harry, III; Holmes-Farley, Stephen Randall; Huval, Chad Cori; Garigapati, Venkata R.; Shackett, Keith K.; Concagh, Danny PATENT ASSIGNEE(S): Geltex Pharmaceuticals, Inc., USA SOURCE: PCT Int. Appl., 104 pp. CODEN: PIXXD2 DOCUMENT TYPE: Patent English LANGUAGE: FAMILY ACC. NUM. COUNT: PATENT INFORMATION: PATENT NO. KIND DATE APPLICATION NO. DATE ----------A1 WO 1999-US15958 19990714 WO 2001005408 20010125 W: AE, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, CA, CH, CN, CU, CZ, DE, DK, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MD, MG, MK, MN, MW, MX, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, UA, UG, UZ, VN, YU, ZA, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM RW: GH, GM, KE, LS, MW, SD, SL, SZ, UG, ZW, AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, BF, BJ, CF, CG, CI, CM, GA, GN, GW, ML, MR, NE, SN, TD, TG AU 9949957 A1 20010205 AU 1999-49957 19990714 PRIORITY APPLN. INFO.: WO 1999-US15958 A 19990714 Polymers having ether and(or) N-contg. side chains are manufd. for use in binding fat for treatment of obesity. A typical polymer was manufd. by radical polymn. of N-decylacrylamide 2.83, 3-acrylamidopropyltrimethylammo nium chloride 18.45, and acrylamide 13.33 g. THERE ARE 10 CITED REFERENCES AVAILABLE FOR REFERENCE COUNT: 10 THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

=> d iall

L18 ANSWER 1 OF 1 CAPLUS COPYRIGHT 2002 ACS

ACCESSION NUMBER:

2001:63835 CAPLUS

DOCUMENT NUMBER:

134:131954

TITLE:

INVENTOR(S):

Fat-binding polymers for use with lipase inhibitors Jozefiak, Thomas Henry; Mandeville, W. Harry, III; Holmes-Farley, Stephen Randall; Huval, Chad Cori; Garigapati, Venkata R.; Shackett, Keith K.; Concagh,

Danny

PATENT ASSIGNEE(S):

Geltex Pharmaceuticals, Inc., USA

SOURCE:

PCT Int. Appl., 104 pp. CODEN: PIXXD2

DOCUMENT TYPE:

Patent

LANGUAGE: English

INT. PATENT CLASSIF.:

MAIN:

A61K031-785

SECONDARY:

A61P003-00; A61K031-785; A61K031-335

CLASSIFICATION:

35-4 (Chemistry of Synthetic High Polymers)

Section cross-reference(s): 63

FAMILY ACC. NUM. COUNT:

PATENT INFORMATION:

	PATENT NO.			KIND DATE			APPLICATION NO. DATE													
				- -						-										
	WO	2001005408			A1		20010125		WO 1999-US15958 19990714											
		W:	ΑE,	AL,	AM,	ΑT,	AU,	ΑZ,	BA,	BB,	BG,	BR,	BY,	CA,	CH,	CN,	CU,	CZ,		
			DE,	DK,	EE,	ES,	FI,	GB,	GD,	GE,	GH,	GM,	HR,	HU,	ID,	IL,	IN,	IS,		
			JP,	KΕ,	KG,	ΚP,	KR,	ΚZ,	LC,	LK,	LR,	LS,	LT,	LU,	LV,	MD,	MG,	MK,		
			MN,	MW,	MX,	NO,	NZ,	PL,	PT,	RO,	RU,	SD,	SE,	SG,	SI,	SK,	SL,	ТJ,		
			TM,	TR,	TT,	UA,	UG,	UZ,	VN,	ΥU,	ZA,	ZW,	AM,	ΑZ,	BY,	KG,	ΚZ,	MD,		
			RU,	ТJ,	TM															
		RW:	GH,	GM,	ΚE,	LS,	MW,	SD,	SL,	SZ,	UG,	ZW,	ΑT,	BE,	CH,	CY,	DE,	DK,		
			ES,	FI,	FR,	GB,	GR,	ΙE,	IT,	LU,	MC,	NL,	PT,	SE,	BF,	ВJ,	CF,	CG,		
			CI,	CM,	GΑ,	GN,	GW,	ML,	MR,	ΝE,	SN,	TD,	TG							
	ΑU	9949	957		A.	1	2001	0205		A	U 19	99-4	9957		1999	0714				
PRIOR	RITY	APP	LN.	INFO	. :				1	WO 1	999-1	US15	958	Α	1999	0714				
		•																		

ABSTRACT:
Polymers having ether and(or) N-contg. side chains are manufd. for use in binding fat for treatment of obesity. A typical polymer was manufd. by

radical polymn. of N-decylacrylamide 2.83, 3-acrylamidopropyltrimethylammonium chloride

18.45, and acrylamide 13.33 g.

SUPPL. TERM: fat binding nitrogen contg side chain polymer manuf;

decylacrylamide acrylaminopropyltrimethylammonium chloride acrylamide copolymer manuf fat binding; ether contg side

chain polymer manuf fat binding

INDEX TERM: Antiobesity agents

Hypertriglyceridemia

(fat-binding polymers for use with lipase inhibitors)

INDEX TERM: Ionene polymers

ROLE: IMF (Industrial manufacture); PRP (Properties); PREP

(Preparation)

(fat-binding polymers for use with lipase inhibitors)

INDEX TERM: Cardo polymers

ROLE: IMF (Industrial manufacture); PRP (Properties); PREP

(Preparation)

(maleimide group-contg. polymers; fat-binding polymers

for use with lipase inhibitors)

INDEX TERM: Quaternary ammonium compounds, preparation

ROLE: IMF (Industrial manufacture); PRP (Properties); PREP

(Preparation)

(polymers; fat-binding polymers for use with lipase

inhibitors)

INDEX TERM: Digestive tract

(steatorrhea; fat-binding polymers for use with lipase

inhibitors)

INDEX TERM: 321904-06-7P, Poly[N,N-diallyl-N-methyl-N-(2,3-

dihydroxypropyl) ammonium chloride] 321904-08-9P

321904-11-4P 321904-13-6P 321904-14-7P

ROLE: IMF (Industrial manufacture); PREP (Preparation) (fat-binding polymers for use with lipase inhibitors)

INDEX TERM: 109-55-7DP, 3-(Dimethylamino)propylamine, reaction products

with ethylene-maleic anhydride alternating copolymer 540-51-2DP, 2-Bromoethanol, reaction products with polyethylenimine 556-52-5DP, Glycidol, reaction products with polyallylamine hydrochloride 590-92-1DP, 3-Bromopropionic acid, reaction products with polyethylenimine 1002-69-3DP, 1-Chlorodecane, reaction products with allylamine-diallyldimethylammonium chloride copolymer and chloroacetic acid 1120-71-4DP, 1,3-Propanesultone, reaction products with polydiallylmethylamine hydrochloride 9002-98-6DP, Polyethylenimine, reaction products with bromopropionic

acid

9039-82-1DP, Polyethylene glycol glycidyl nonylphenyl

ether,

reaction products with polydiallylmethylamine hydrochloride 25805-17-8DP, Poly(2-ethyl-2-oxazoline), partially 26063-69-4DP, Polydiallylamine hydrochloride, hydrolyzed functionalized 26403-72-5DP, Polyethylene glycol diglycidyl ether, reaction products with polydiallylmethylamine hydrochloride 26427-01-0P. Poly(3-acrylamidopropyltrimethylammonium chloride) 32765-81-4DP, 6-Bromohexyltrimethylammonium bromide, reaction products with polydiallylamine hydrochloride 34447-60-4P, Acrylamide-diallylammonium chloride copolymer 40349-67-5DP, Polyethylene glycol glycidyl methyl ether, reaction products with polydiallylmethylamine hydrochloride 51729-06-7P, Diallyldimethylammonium chloride-vinyl alcohol copolymer 53694-17-0P, Acrylic aciddiallyldimethylammonium chloride copolymer 55553-13-4DP, Poly(diallylmethylamine), functionalized 62238-80-6DP, Polydiallylamine, functionalized 68240-11-9P, Acrylamide-diallylmethylamine hydrochloride copolymer 71550-12-4DP, Polyallylamine hydrochloride, functionalized 73354-75-3P, Poly(N,N-diallyl-2-hydroxyethylamine 75150-29-7P, Acrylamide-(3hydrochloride) acrylamidopropyl) trimethylammonium chloride copolymer 76123-63-2P 83601-65-4P,

(3-Acrylamidopropyl)trimethylammo

nium chloride-styrene copolymer 84154-72-3P, Acrylamide-N-[3-(dimethylamino)propyl]acrylamide copolymer 86630-59-3DP, Polyethylene glycol glycidyl lauryl ether, reaction products with polydiallylmethylamine hydrochloride 106973-21-1DP, Ethylene-maleic anhydride alternating copolymer, reaction products with dimethylaminopropylamine 131479-66-8P, (3-Acrylamidopropyl) trimethylammonium chloride-acrylic acid copolymer 151274-11-2P, (3-Acrylamidopropyl)trimethylammonium chloride-N-vinyl-2pyrrolidone copolymer 164719-55-5DP, Allylaminediallyldimethylammonium chloride copolymer, reaction products with chloroacetic acid 165957-71-1P, Acrylamide-3-methyl-1-vinylimidazolium chloride copolymer 321903-78-0P,

Acrylamide-(3-acrylamidopropyl)trimethylammoni

um chloride-N-decylacrylamide copolymer 321903-79-1P, Acrylamide-(3-acrylamidopropyl) trimethylammonium chloride-N,N-didecylacrylamide copolymer 321903-80-4P, Acrylamide-(3-acrylamidopropyl) trimethylammonium chloride-N-phenylacrylamide copolymer 321903-81-5P, Acrylamide-(3-acrylamidopropyl) trimethylammonium chloride-N-benzylacrylamide copolymer 321903-82-6P, (3-Acrylamidopropyl) trimethylammonium chloride-N-tert-

7

octylacrylamide copolymer 321903-83-7P, (3-Acrylamidopropyl) trimethylammonium chloride-N-321903-85-9P, butylacrylamide copolymer Poly(2-methacryloyloxyethyl-tert-butylamine hydrochloride) 321903-86-0P 321903-87-1P 321903-88-2P, Acrylamide-(3-acrylamidopropyl)trimethylammonium chloride-N-octadecylacrylamide copolymer Acrylamide-(3-acrylamidopropyl) trimethylammonium chloride-N-methyl-N-octadecylacrylamide copolymer 321903-91-7P, Acrylamide-N-dodecylacrylamide-3-methyl-1vinylimidazolium chloride copolymer 321903-92-8P, (3-Acrylamidopropyl) trimethylammonium chloride-Nethylacrylamide copolymer 321903-93-9P, (3-Acrylamidopropyl)trimethylammonium chloride-polyethylene glycol acrylate methyl ether graft copolymer 321903-94-0P 321903-95-1P 321903-96-2P 321903-97-3P 321903-98-4P, Acrylamide-N-[3-(dimethylamino)propyl]acrylamide-Ndodecylacrylamide copolymer 321904-00-1P 321904-01-2P, Diallyldimethylammonium chloride-polyethylene glycol acrylate methyl ether graft copolymer 321904-02-3P 321904-03-4P, Acrylamide-(3-acrylamidopropyl)trimethylammoni um chloride-N-octylacrylamide copolymer 321904-04-5P, Acrylamide-(3-acrylamidopropyl)trimethylammonium chloride-methylenebisacrylamide-N-dodecylacrylamide copolymer 321904-05-6P 321904-16-9P 321936-94-1P, (3-Acrylamidopropyl)trimethylammonium chloride-ethylene oxide graft copolymer methyl ether 321936-96-3P, Diallyldimethylammonium chloride-propylene oxide graft copolymer methyl ether 321936-97-4P, Diallyldimethylammonium chloride-polypropylene glycol acrylate methyl ether graft copolymer 321936-99-6P, Diallyldimethylammonium chloride-ethylene oxide graft copolymer methyl ether ROLE: IMF (Industrial manufacture); PRP (Properties); PREP (Preparation) (fat-binding polymers for use with lipase inhibitors) INDEX TERM: 26063-69-4P, Polydiallylamine hydrochloride 29566-78-7P, Poly (N, N-diallylmethylamine hydrochloride) ROLE: IMF (Industrial manufacture); PRP (Properties); RCT (Reactant); PREP (Preparation) (fat-binding polymers for use with lipase inhibitors) INDEX TERM: 26062-79-3, Polydiallyldimethylammonium chloride 321903-99-5, Bis(2-chloroethyl) 26658-46-8 ether-1,3-bis[3-(dimethylamino)propyl]urea alternating copolymer ROLE: PRP (Properties) (fat-binding polymers for use with lipase inhibitors) INDEX TERM: 74-88-4, Methyl iodide, reactions 1072-63-5, 1-Vinylimidazole ROLE: RCT (Reactant) (monomer precursor; fat-binding polymers for use with lipase inhibitors) INDEX TERM: 13474-25-4P, 3-Methyl-1-vinylimidazolium chloride 32171-39-4P, Polyethylene glycol methyl ether acrylate ROLE: IMF (Industrial manufacture); RCT (Reactant); PREP (Preparation) (monomer; fat-binding polymers for use with lipase inhibitors) REFERENCE COUNT: 10 THERE ARE 10 CITED REFERENCES AVAILABLE FOR THIS

RECORD.

REFERENCE(S):

- (1) Day, C; US 5900233 A 1999 CAPLUS
- (2) Fields, J; US 4211765 A 1980 CAPLUS
- (3) Hadvary, P; US 4598089 A 1986 CAPLUS
- (4) Hoffmann La Roche; EP 0129748 A 1985 CAPLUS
- (5) Holmes-Farley, S; US 5607669 A 1997 CAPLUS
- (6) Holmes-Farley, S; US 5618530 A 1997 CAPLUS
- (7) Holmes-Farley, S; US 5679717 A 1997 CAPLUS
- (8) Holmes-Farley, S; US 5900475 A 1999 CAPLUS
- (9) Page, J; US 4432968 A 1984 CAPLUS
- (10) SjOstrOm, L; LANCET 1998, V352(9123), P167 CAPLUS

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Calculated physical property data is now available. See HELP PROPERTIES for more information. See STNote 27, Searching Properties in the CAS Registry File, for complete details: http://www.cas.org/ONLINE/STN/STNOTES/stnotes27.pdf

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=> STR 26063-69-4

26063-69-4 MAY NOT BE USED AS A MODEL COMPONENTS

6147-66-6 C6 H11 N ?H Cl Single Atom Fragment 1 Cl

ENTER NAME OF STRUCTURE TO BE RECALLED (NONE): END

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THIS FEATURE IS NOT AVAILABLE FOR THE SELECTED CAS RN

=> s 26063-69-4P

L19 0 26063-69-4P

(26063-69-4P/RN)

=> s 26063-69-4

L20 1 26063-69-4

(26063-69-4/RN)

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NEWS 5 Feb 19 Access via Tymnet and SprintNet Eliminated Effective 3/31/02

NEWS 6 Mar 08 Gene Names now available in BIOSIS

NEWS 7 Mar 22 TOXLIT no longer available

NEWS 8 Mar 22 TRCTHERMO no longer available

NEWS 9 Mar 28 US Provisional Priorities searched with P in CA/CAplus and USPATFULL

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NEWS 15 Apr 19 US Patent Applications available in IFICDB, IFIPAT, and IFIUDB

NEWS 16 Apr 22 Paccade from IP com available in CAPLUS HCAPLUS and

NEWS 16 Apr 22 Records from IP.com available in CAPLUS, HCAPLUS, and ZCAPLUS

NEWS 17 Apr 22 BIOSIS Gene Names now available in TOXCENTER

NEWS 18 Apr 22 Federal Research in Progress (FEDRIP) now available

NEWS 19 May 31 PCTFULL to be reloaded. File temporarily unavailable.

NEWS 20 Jun 03 New e-mail delivery for search results now available

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CURRENT MACINTOSH VERSION IS V6.0a(ENG) AND V6.0Ja(JP),

AND CURRENT DISCOVER FILE IS DATED 05 FEBRUARY 2002

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^{=&}gt; s diallyl?ammonium

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Symbol	Definition	Example	Retrieves
?	Any number of characters	S STEROID?	STEROID STEROIDS STEROIDAL
	Left truncation in some fields	S ?METHANE	METHANE TRIHALOMETHANE NITROMETHANE
#	One or zero characters at the end of a term	S VAPOR#	VAPOR VAPORS
!	One character at the end of or within a term	S WOM!N	WOMAN WOMEN
!#	At least one character, and no more than two, at the end of a term	S STEROID!#	STEROIDS STEROIDAL

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for some fields in certain files. Left truncation of a search term can be used alone or, in some cases, in combination with right truncation. The cases which may not be used are: '?text#', '#text?' and '!text?'. To learn whether any fields in the current file have left truncation, enter HELP SFIELDS at an arrow prompt (=>) in the file.

=> s allylammonium

L1 145 ALLYLAMMONIUM

=> s diallylammonium

L2 85 DIALLYLAMMONIUM

=> s 12 and dihydroxylprop?

0 DIHYDROXYLPROP?

L3 0 L2 AND DIHYDROXYLPROP?

=> s 12 and dihydroxyprop?

15207 DIHYDROXYPROP?

L4 0 L2 AND DIHYDROXYPROP?

=> log y

COST IN U.S. DOLLARS

SINCE FILE TOTAL ENTRY SESSION

FULL ESTIMATED COST 19.80 20.01

STN INTERNATIONAL LOGOFF AT 14:11:33 ON 04 JUN 2002

L21 ANSWER 11 OF 15 CAPLUS COPYRIGHT 2002 ACS DUPLICATE 1

ACCESSION NUMBER: 1999:368919 CAPLUS

DOCUMENT NUMBER: 131:139277

TITLE: Structural and ionic determinants of

5-nitro-2-(3-phenylpropylamino)-benzoic acid block of

the CFTR chloride channel

AUTHOR(S): Walsh, Kenneth B.; Long, Kathryn J.; Shen, Xufeng

CORPORATE SOURCE: Department of Pharmacology, School of Medicine,

University of South Carolina, Columbia, SC, 29208,

USA

SOURCE: British Journal of Pharmacology (1999), 127(2),

369-376

CODEN: BJPCBM; ISSN: 0007-1188

PUBLISHER:

Stockton Press

DOCUMENT TYPE:

Journal

LANGUAGE:

English

AB The goals of this study were to identify the structural components required for arylaminobenzoate block of the **cystic**

fibrosis transmembrane conductance regulator (CFTR) chloride channel and to det. the involvement of two pos. charged amino acid

residues, found within the channel, in drug binding. Wild-type and mutant

CFTR chloride channels were expressed in Xenopus oocytes and CFTR currents

measured using the two microelectrode voltage clamp. Block of the wild-type CFTR current by 5-nitro-2-(3-phenylpropylamino)-benzoate (NPPB) occurred in a voltage-dependent manner with preferential inhibition of

inward currents (Kd=166 .mu.M at -90 mV). Removal of the Ph ring from the $\,$

aliph. chain of NPPB, with the compd. 2-butylamino-5-nitrobenzoic acid, caused only a small change in CFTR inhibition (Kd=243 .mu.M), while addn. of an extra Ph ring at this position

(5-nitro-2-(3,3-diphenylpropylamino)-

benzoic acid) increased drug potency (Kd=58 .mu.M). In contrast, removal of the benzoate ring (2-amino-4-phenylbutyric acid) or the 5-nitro group (2-(3-phenylpropylamino)-benzoic acid) of NPPB severely limited drug block of the wild-type channel. NPPB inhibition of CFTR currents in oocytes expressing the mutants K335E and R347E also occurred in a voltage-dependent manner. However, the Kds for NPPB block were increased to 371 and 1573 .mu.M, for the K335E and R347E mutants, resp. NPPB block of the inward wild-type CFTR current was reduced in the presence of 10 mM of the permeant anion SCN-. These studies present the first step in the development of high affinity probes to the CFTR channel.

REFERENCE COUNT:

26 THERE ARE 26 CITED REFERENCES AVAILABLE FOR

THIS

the

RECORD. ALL CITATIONS AVAILABLE IN THE RE

FORMAT

L21 ANSWER 10 OF 15 MEDLINE

ACCESSION NUMBER: 1999447279 MEDLINE

DOCUMENT NUMBER: 99447279 PubMed ID: 10516210

TITLE: PBA increases CFTR expression but at high doses inhibits

Cl(-) secretion in Calu-3 airway epithelial cells.

AUTHOR: Loffing J; Moyer B D; Reynolds D; Stanton B A

CORPORATE SOURCE: Department of Physiology, Dartmouth Medical School,

Hanover, New Hampshire 03755, USA.

CONTRACT NUMBER: CA-23108 (NCI)

DK-45881 (NIDDK) HL-45881 (NHLBI)

SOURCE: AMERICAN JOURNAL OF PHYSIOLOGY, (1999 Oct) 277 (4 Pt 1)

L700-8.

Journal code: 3U8; 0370511. ISSN: 0002-9513.

PUB. COUNTRY: United States

Journal; Article; (JOURNAL ARTICLE)

LANGUAGE: English

FILE SEGMENT: Priority Journals

ENTRY MONTH: 199911

ENTRY DATE: Entered STN: 20000111

Last Updated on STN: 20000111 Entered Medline: 19991122

AB Sodium 4-phenylbutyrate (PBA), a short-chain fatty acid, has been approved

to treat patients with urea cycle enzyme deficiencies and is being evaluated in the management of sickle cell disease, thalassemia, cancer, and cystic fibrosis (CF). Because relatively little is known about the effects of PBA on the expression and function of the wild-type CF transmembrane conductance regulator (wt CFTR), the goal of this study was to examine the effects of PBA and related compounds on wt CFTR-mediated Cl(-) secretion. To this end, we studied Calu-3 cells, a human airway cell line that expresses endogenous wt CFTR and has a serous cell phenotype. We report that chronic treatment of Calu-3 cells with a high concentration (5 mM) of PBA, sodium butyrate, or sodium valproate

but

not of sodium acetate reduced basal and 8-(4-chlorophenylthio)-cAMP-stimulated Cl(-) secretion. Paradoxically, PBA enhanced CFTR protein expression 6- to 10-fold and increased the intensity of CFTR staining in the apical plasma membrane. PBA also increased protein expression of Na(+)-K(+)-ATPase. PBA reduced CFTR Cl(-) currents across the apical membrane but had no effect on Na(+)-K(+)-ATPase activity in the basolateral membrane. Thus a high concentration of PBA (5 mM) reduces Cl(-) secretion by inhibiting CFTR Cl(-) currents across the apical membrane. In contrast, lower therapeutic concentrations of PBA (0.05-2

mM)

had no effect on cAMP-stimulated Cl(-) secretion across Calu-3 cells. We conclude that PBA concentrations in the therapeutic range are unlikely to have a negative effect on Cl(-) secretion. However, concentrations >5 mM might reduce transepithelial Cl(-) secretion by serous cells in submucosal

glands in individuals expressing wt CFTR.

L10 ANSWER 40 OF 40 MEDLINE

ACCESSION NUMBER: 81109797 MEDLINE

DOCUMENT NUMBER: 81109797 PubMed ID: 7193034

TITLE: The bile acid binding and hypocholesterolemic action of

two

water-soluble polymers.

AUTHOR: Kuron G W; Grier N; Huff J W

SOURCE: ATHEROSCLEROSIS, (1980 Nov) 37 (3) 353-60.

Journal code: 95X; 0242543. ISSN: 0021-9150.

PUB. COUNTRY: Netherlands

Journal; Article; (JOURNAL ARTICLE)

LANGUAGE: English

FILE SEGMENT: Priority Journals

ENTRY MONTH: 198103

ENTRY DATE: Entered STN: 19900316

Last Updated on STN: 19900316 Entered Medline: 19810317

AB The in vitro bile acid binding properties of 2 water-soluble, linear, cationic resins, poly-[(dimethylimino)trimethylene chloride] or 3,3-ione C1, and poly-diallyldimethylammonium chloride) or CAT-FLOC were determined. Both polymers were substantially more active than cholestyramine. All were compared for hypocholesterolemic effect in normo-cholesterolemic dogs. CAT-FLOC and 3,3-ionene C1, administered at 1.8 and 1.2 g/day, respectively, exhibited cholesterol-lowering action equivalent to cholesteryramine given at 12 g/day. The results of this study suggest that effective reduction of plasma cholesterol may be achieved with significantly lower doses of bile acid sequestrants.

L10 ANSWER 39 OF 40 MEDLINE

ACCESSION NUMBER: 82061521 MEDLINE

DOCUMENT NUMBER: 82061521 PubMed ID: 7302387

TITLE: Effect of three bile acid binding polymers on the

biosynthesis of 14C-cholesterol from 14C-sodium acetate in

the rat

AUTHOR: Gilfillan J L; Huff J W

SOURCE: RESEARCH COMMUNICATIONS IN CHEMICAL PATHOLOGY AND

PHARMACOLOGY, (1981 Aug) 33 (2) 373-6.

Journal code: R62; 0244734. ISSN: 0034-5164.

PUB. COUNTRY: United States

Journal; Article; (JOURNAL ARTICLE)

LANGUAGE: English

FILE SEGMENT: Priority Journals

ENTRY MONTH: 198201

ENTRY DATE: Entered STN: 19900316

Last Updated on STN: 19900316 Entered Medline: 19820109

AB The relative activity of three bile acid binding polymers in increasing cholesterol biosynthesis in the rat from 14C-acetate was determined by measuring blood levels of 14C-cholesterol after intraperitoneally administered 14C-acetate. CAT-FLOC and 3,3-ionene were 4-5 times more active than cholestyramine in this study which correlated well with the results of hypocholesteremic testing in dogs.

L10 ANSWER 36 OF 40 MEDLINE

ACCESSION NUMBER: 96091252 MEDLINE

DOCUMENT NUMBER: 96091252 PubMed ID: 7489112

TITLE: [The antimutagenic activity of ternary diallyl

copolymers].

Antimutagennaia aktivnost' troinykh sopolimerov

diallil'nogo riada.

AUTHOR: Aleksandrova V A; Kotliarova E B; Odin A P; Domnina N S;

Shevchenko V A; Topchiev D A

SOURCE: RADIATSIONNAIA BIOLOGIIA, RADIOECOLOGIIA, (1995 Sep-Oct)

35

(5) 746-51.

Journal code: BWZ; 9317212. ISSN: 0869-8031.

PUB. COUNTRY: RUSSIA: Russian Federation

Journal; Article; (JOURNAL ARTICLE)

LANGUAGE: Russian

FILE SEGMENT: Priority Journals

ENTRY MONTH: 199601

ENTRY DATE: Entered STN: 19960125

Last Updated on STN: 19960125 Entered Medline: 19960102

AB Antimutagenic activity of triple copolymers of diallyl origin was investigated by animal cell test (mouse bone marrow erythrocytes, 1.5 Gy of gamma irradiation) and by plant cell test (seeds of barley, 5 Gy of gamma irradiation). Effective protection of genetic structure was achieved

owing to combination of moderate antimutagenic activity of the polymer matrix and scavenging ability of sterically hindered phenols in the polymer side chain.

L10 ANSWER 33 OF 40 CAPLUS COPYRIGHT 2002 ACS

ACCESSION NUMBER: 1985:561469 CAPLUS

DOCUMENT NUMBER: 103:161469

TITLE: Microcapsules with permeable or semipermeable walls

and a liquid core

INVENTOR(S): Loth, Fritz; Dautzenberg, Horst; Pommerening, Klaus PATENT ASSIGNEE(S): Akademie der Wissenschaften der DDR, Ger. Dem. Rep.

SOURCE: Ger. (East), 12 pp. Addn. to Ger. (East) 160,393.

CODEN: GEXXA8

DOCUMENT TYPE: Patent LANGUAGE: German

FAMILY ACC. NUM. COUNT: 2

PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
	- -			
DD 218734	A4	19850213	DD 1981-232617	19810817
DD 160393	T	19830727	DD 1980-225200	19801114
PRIORITY APPLN. INFO.:			DD 1980-225200	19801114
3D	<i>-</i>	1 1		

AB Capsule walls are formed by pptn. of anionic and cationic polyelectrolytes

at their interface. The anionic polyelectrolytes are sulfate or carboxylate contg. polysaccharide and/or synthetic polymers, and the cationic polyelectrolytes include quaternary ammonium surfactants and/or dyes. The microcapsules can be used for sepn. processes in preparative and anal. chem. and biochem., and in **pharmacy**, **medicine**, and agrochem. and food industries. Thus, 0.2 g Na cellulose sulfate

[9005-22-5] with a degree of substitution of 0.4 was dissolved in 9.8 g H2O, and the soln. was pressed through a 0.2-mm inner diam. capillary and dropped from a height of 30 cm into a stirred bath contg. 1% aq.

methylene

blue [61-73-4]. After 30 min the capsules formed were decanted and washed with H2O. The deep-blue capsules had a diam. of 3-5 mm.

L6 ANSWER 2 OF 2 CAPLUS COPYRIGHT 2002 ACS

ACCESSION NUMBER: 1993:256861 CAPLUS

DOCUMENT NUMBER: 118:256861

TITLE: Rosin emulsion sizes and dispersing agents for

improving their stability and endurance to water

hardness and acidity stress

INVENTOR(S): Niike, Hitoshi; Sakuraba, Noriko

PATENT ASSIGNEE(S): Daiichi Kogyo Seiyaku Co., Ltd., Japan

SOURCE: Jpn. Kokai Tokkyo Koho, 17 pp.

CODEN: JKXXAF

DOCUMENT TYPE: Patent

LANGUAGE: Japanese FAMILY ACC. NUM. COUNT: 1

PATENT INFORMATION:

PATENT NO. KIND DATE APPLICATION NO. DATE

JP 04333694 A2 19921120 JP 1991-128646 19910430

AB The title agents are copolymers and/or their salts prepd. from (meth) acrylate esters and/or styrene compds., and comonomers bearing amino and/or ammonium groups in the presence of 2,4-diphenyl-4-methyl-1-pentene (I) chain-transfer agents. Thus, 2,2'-azobis(2,4-dimethylvaleronitrile)-initiated polymn. of Et acrylate, Me methacrylate, sec-Bu acrylate, [(methacryloyloxy)ethyl]trimethylammonium chloride, and [(methacryloyloxy)-2-hydroxypropyl]trimethylammonium chloride in the presence of I gave a copolymer with mol. wt. 25,000, which was used to disperse a fumaric acid-fortified rosin into a stable emulsion.

L18 ANSWER 1 OF 1 CAPLUS COPYRIGHT 2002 ACS ACCESSION NUMBER: 2001:63835 CAPLUS

DOCUMENT NUMBER: 134:131954

TITLE: Fat-binding polymers for use with lipase inhibitors INVENTOR (S): Jozefiak, Thomas Henry; Mandeville, W. Harry, III; Holmes-Farley, Stephen Randall; Huval, Chad Cori; Garigapati, Venkata R.; Shackett, Keith K.; Concagh,

Danny

PATENT ASSIGNEE(S): Geltex Pharmaceuticals, Inc., USA

SOURCE:

PCT Int. Appl., 104 pp.

CODEN: PIXXD2

DOCUMENT TYPE: Patent LANGUAGE: English

INT. PATENT CLASSIF.:

MAIN: A61K031-785

SECONDARY: A61P003-00; A61K031-785; A61K031-335

CLASSIFICATION: 35-4 (Chemistry of Synthetic High Polymers)

Section cross-reference(s): 63

FAMILY ACC. NUM. COUNT:

PATENT INFORMATION:

PATENT NO. KIND DATE APPLICATION NO. DATE ---------WO 2001005408 A1 20010125 WO 1999-US15958 19990714 W: AE, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, CA, CH, CN, CU, CZ, DE, DK, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MD, MG, MK, MN, MW, MX, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, UA, UG, UZ, VN, YU, ZA, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM RW: GH, GM, KE, LS, MW, SD, SL, SZ, UG, ZW, AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, BF, BJ, CF, CG, CI, CM, GA, GN, GW, ML, MR, NE, SN, TD, TG AU 9949957 A1 20010205 AU 1999-49957 19990714

PRIORITY APPLN. INFO.: WO 1999-US15958 A 19990714

Polymers having ether and(or) N-contg. side chains are manufd. for use in binding fat for treatment of obesity. A typical polymer was manufd. by

polymn. of N-decylacrylamide 2.83, 3-acrylamidopropyltrimethylammonium chloride

18.45, and acrylamide 13.33 g.

SUPPL. TERM: fat binding nitrogen contg side chain polymer manuf;

decylacrylamide acrylaminopropyltrimethylammonium chloride acrylamide copolymer manuf fat binding; ether contg side

chain polymer manuf fat binding

INDEX TERM: Antiobesity agents

Hypertriglyceridemia

(fat-binding polymers for use with lipase inhibitors)

INDEX TERM: Ionene polymers

ROLE: IMF (Industrial manufacture); PRP (Properties); PREP

(Preparation)

(fat-binding polymers for use with lipase inhibitors)

INDEX TERM: Cardo polymers

ROLE: IMF (Industrial manufacture); PRP (Properties); PREP

(Preparation)

(maleimide group-contg. polymers; fat-binding polymers

for use with lipase inhibitors)

INDEX TERM:

Quaternary ammonium compounds, preparation

ROLE: IMF (Industrial manufacture); PRP (Properties); PREP

(Preparation)

(polymers; fat-binding polymers for use with lipase

inhibitors)

INDEX TERM:

Digestive tract

(steatorrhea; fat-binding polymers for use with lipase

inhibitors)

INDEX TERM:

321904-06-7P, Poly[N,N-diallyl-N-methyl-N-(2,3-dihydroxypropyl)ammonium chloride] 321904-08-9P

321904-11-4P 321904-13-6P 321904-14-7P

ROLE: IMF (Industrial manufacture); PREP (Preparation) (fat-binding polymers for use with lipase inhibitors)

INDEX TERM:

109-55-7DP, 3-(Dimethylamino) propylamine, reaction products with ethylene-maleic anhydride alternating copolymer 540-51-2DP, 2-Bromoethanol, reaction products with

polyethylenimine 556-52-5DP, Glycidol, reaction products

with polyallylamine hydrochloride 590-92-1DP, 3-Bromopropionic acid, reaction products with

polyethylenimine 1002-69-3DP, 1-Chlorodecane, reaction products with allylamine-diallyldimethylammonium chloride

copolymer and chloroacetic acid 1120-71-4DP, 1,3-Propanesultone, reaction products with

polydiallylmethylamine hydrochloride 9002-98-6DP, Polyethylenimine, reaction products with bromopropionic

acid

9039-82-1DP, Polyethylene glycol glycidyl nonylphenyl

ether,

reaction products with polydiallylmethylamine hydrochloride 25805-17-8DP, Poly(2-ethyl-2-oxazoline), partially 26063-69-4DP, Polydiallylamine hydrochloride, hydrolyzed functionalized 26403-72-5DP, Polyethylene glycol diglycidyl ether, reaction products with polydiallylmethylamine hydrochloride 26427-01-0P. Poly(3-acrylamidopropyltrimethylammonium chloride) 32765-81-4DP, 6-Bromohexyltrimethylammonium bromide, reaction products with polydiallylamine hydrochloride 34447-60-4P, Acrylamide-diallylammonium chloride copolymer 40349-67-5DP, Polyethylene glycol glycidyl methyl ether, reaction products with polydiallylmethylamine hydrochloride 51729-06-7P, Diallyldimethylammonium chloride-vinyl alcohol copolymer 53694-17-0P, Acrylic aciddiallyldimethylammonium chloride copolymer 55553-13-4DP, Poly(diallylmethylamine), functionalized 62238-80-6DP, Polydiallylamine, functionalized 68240-11-9P, Acrylamide-diallylmethylamine hydrochloride copolymer 71550-12-4DP, Polyallylamine hydrochloride, functionalized 73354-75-3P, Poly(N,N-diallyl-2-hydroxyethylamine hydrochloride) 75150-29-7P, Acrylamide-(3acrylamidopropyl)trimethylammonium chloride copolymer

(3-Acrylamidopropyl)trimethylammo

76123-63-2P

nium chloride-styrene copolymer 84154-72-3P, Acrylamide-N-[3-(dimethylamino)propyl]acrylamide copolymer 86630-59-3DP, Polyethylene glycol glycidyl lauryl ether, reaction products with polydiallylmethylamine hydrochloride 106973-21-1DP, Ethylene-maleic anhydride alternating copolymer, reaction products with dimethylaminopropylamine 131479-66-8P, (3-Acrylamidopropyl)trimethylammonium chloride-acrylic acid copolymer 151274-11-2P,

83601-65-4P,

aciu

echer,

. - -

(3-Acrylamidopropyl)trimethylammonium chloride-N-vinyl-2-pyrrolidone copolymer 164719-55-5DP, Allylamine-diallyldimethylammonium chloride copolymer, reaction products with chloroacetic acid 165957-71-1P, Acrylamide-3-methyl-1-vinylimidazolium chloride copolymer 321903-78-0P, Acrylamidopropyl)trimethylammoni um chloride-N-decylacrylamide copolymer 321903-79-1P, Acrylamide-(3-acrylamidopropyl)trimethylammonium chloride-N,N-didecylacrylamide copolymer 321903-80-4P,

Acrylamide-(3-acrylamidopropyl)trimethylammonium chloride-N-phenylacrylamide copolymer 321903-81-5P, Acrylamide-(3-acrylamidopropyl)trimethylammonium chloride-N-benzylacrylamide copolymer 321903-82-6P, (3-Acrylamidopropyl) trimethylammonium chloride-N-tertoctylacrylamide copolymer 321903-83-7P, (3-Acrylamidopropyl)trimethylammonium chloride-Nbutylacrylamide copolymer 321903-85-9P, Poly(2-methacryloyloxyethyl-tert-butylamine hydrochloride) 321903-86-0P 321903-87-1P 321903-88-2P. Acrylamide - (3-acrylamidopropyl) trimethylammonium chloride-N-octadecylacrylamide copolymer 321903-89-3P, Acrylamide-(3-acrylamidopropyl)trimethylammonium chloride-N-methyl-N-octadecylacrylamide copolymer 321903-91-7P, Acrylamide-N-dodecylacrylamide-3-methyl-1vinylimidazolium chloride copolymer 321903-92-8P, (3-Acrylamidopropyl) trimethylammonium chloride-Nethylacrylamide copolymer 321903-93-9P, (3-Acrylamidopropyl)trimethylammonium chloride-polyethylene glycol acrylate methyl ether graft copolymer 321903-94-0P 321903-98-4P, 321903-95-1P 321903-96-2P 321903-97-3P Acrylamide-N-[3-(dimethylamino)propyl]acrylamide-Ndodecylacrylamide copolymer 321904-00-1P 321904-01-2P, Diallyldimethylammonium chloride-polyethylene glycol acrylate methyl ether graft copolymer 321904-02-3P 321904-03-4P,

Acrylamide-(3-acrylamidopropyl)trimethylammoni

um chloride-N-octylacrylamide copolymer 321904-04-5P, Acrylamide-(3-acrylamidopropyl)trimethylammonium chloride-methylenebisacrylamide-N-dodecylacrylamide copolymer 321904-05-6P 321904-16-9P 321936-94-1P, (3-Acrylamidopropyl)trimethylammonium chloride-ethylene oxide graft copolymer methyl ether 321936-96-3P, Diallyldimethylammonium chloride-propylene oxide graft copolymer methyl ether 321936-97-4P, Diallyldimethylammonium chloride-polypropylene glycol acrylate methyl ether graft copolymer 321936-99-6P, Diallyldimethylammonium chloride-ethylene oxide graft copolymer methyl ether ROLE: IMF (Industrial manufacture); PRP (Properties); PREP (Preparation)

INDEX TERM:

(fat-binding polymers for use with lipase inhibitors) 26063-69-4P, Polydiallylamine hydrochloride 29566-78-7P, Poly(N,N-diallylmethylamine hydrochloride) ROLE: IMF (Industrial manufacture); PRP (Properties); RCT (Reactant); PREP (Preparation)

INDEX TERM:

(fat-binding polymers for use with lipase inhibitors) 26062-79-3, Polydiallyldimethylammonium chloride 26658-46-8 321903-99-5, Bis(2-chloroethyl) ether-1,3-bis[3-(dimethylamino)propyl]urea alternating

copolymer

ROLE: PRP (Properties)

(fat-binding polymers for use with lipase inhibitors)

74-88-4, Methyl iodide, reactions 1072-63-5,

1-Vinylimidazole

ROLE: RCT (Reactant)
(monomer precursor; fat-binding polymers for use with

lipase inhibitors)

INDEX TERM: 13474-25-4P, 3-Methyl-1-vinylimidazolium chloride

32171-39-4P, Polyethylene glycol methyl ether acrylate ROLE: IMF (Industrial manufacture); RCT (Reactant); PREP

(Preparation)

10

(monomer; fat-binding polymers for use with lipase

inhibitors)
THERE ARE 10 CITED REFERENCES AVAILABLE FOR THIS

RECORD.
REFERENCE(S): (1) Day, C; US 5900233 A 1999 CAPLUS

(1) Day, C; US 5900233 A 1999 CAPLUS (2) Fields, J; US 4211765 A 1980 CAPLUS

(3) Hadvary, P; US 4598089 A 1986 CAPLUS

(4) Hoffmann La Roche; EP 0129748 A 1985 CAPLUS

(5) Holmes-Farley, S; US 5607669 A 1997 CAPLUS

(6) Holmes-Farley, S; US 5618530 A 1997 CAPLUS

(7) Holmes-Farley, S; US 5679717 A 1997 CAPLUS

(8) Holmes-Farley, S; US 5900475 A 1999 CAPLUS

(9) Page, J; US 4432968 A 1984 CAPLUS

(10) SjOstrOm, L; LANCET 1998, V352(9123), P167 CAPLUS

=> FILE REG

INDEX TERM:

REFERENCE COUNT:

Trying 3106016892...Open

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LOGINID:ssspta1617sxw
PASSWORD:

TERMINAL (ENTER 1, 2, 3, OR ?):2

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Welcome to STN International
NEWS
                Web Page URLs for STN Seminar Schedule - N. America
NEWS
                IMSworld Pharmaceutical Company Directory name change
        Sep 17
                to PHARMASEARCH
NEWS
     3 Oct 09
                Korean abstracts now included in Derwent World Patents
                Index
     4 Oct 09
NEWS
                Number of Derwent World Patents Index updates increased
NEWS 5 Oct 15
                Calculated properties now in the REGISTRY/ZREGISTRY File
NEWS 6 Oct 22
                Over 1 million reactions added to CASREACT
NEWS 7 Oct 22
                DGENE GETSIM has been improved
NEWS 8 Oct 29
                AAASD no longer available
NEWS 9 Nov 19 New Search Capabilities USPATFULL and USPAT2
NEWS 10 Nov 19
                TOXCENTER(SM) - new toxicology file now available on STN
NEWS 11 Nov 29
                COPPERLIT now available on STN
NEWS 12 Nov 29 DWPI revisions to NTIS and US Provisional Numbers
NEWS 13 Nov 30 Files VETU and VETB to have open access
NEWS 14 Dec 10 WPINDEX/WPIDS/WPIX New and Revised Manual Codes for 2002
NEWS 15 Dec 10 DGENE BLAST Homology Search
NEWS 16 Dec 17 WELDASEARCH now available on STN
NEWS 17 Dec 17 STANDARDS now available on STN
NEWS 18 Dec 17 New fields for DPCI
NEWS 19 Dec 19
                CAS Roles modified
NEWS 20 Dec 19
                1907-1946 data and page images added to CA and CAplus
NEWS 21
        Jan 25
                BLAST(R) searching in REGISTRY available in STN on the Web
NEWS 22
        Jan 25
                Searching with the P indicator for Preparations
NEWS 23
        Jan 29
                FSTA has been reloaded and moves to weekly updates
NEWS 24
        Feb 01
                DKILIT now produced by FIZ Karlsruhe and has a new update
                frequency
NEWS 25
        Feb 19
                Access via Tymnet and SprintNet Eliminated Effective 3/31/02
NEWS EXPRESS
             February 1 CURRENT WINDOWS VERSION IS V6.0d,
             CURRENT MACINTOSH VERSION IS V6.0a(ENG) AND V6.0Ja(JP),
             AND CURRENT DISCOVER FILE IS DATED 05 FEBRUARY 2002
NEWS HOURS
             STN Operating Hours Plus Help Desk Availability
NEWS INTER
             General Internet Information
NEWS LOGIN
             Welcome Banner and News Items
NEWS PHONE
             Direct Dial and Telecommunication Network Access to STN
NEWS WWW
             CAS World Wide Web Site (general information)
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